

STIC-Biotech/ChemLib

93134

From: Stucker, Jeffrey  
Sent: Monday, May 05, 2003 10:36 AM  
To: STIC-Biotech/ChemLib  
Subject: 09/868399

Please search SEQ ID NO 1. The peptide is limited to no more than 60 amino acids in the claims.

Thanks,  
Jeff Stucker  
1648  
308-4237  
mail: 8E12

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STIC/CHEM. DIVISION  
(STIC)

Searcher: \_\_\_\_\_  
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TYPE OF SEARCH:

NA Sequences: \_\_\_\_\_  
AA Sequences: \_\_\_\_\_  
Structures: \_\_\_\_\_  
Bibliographic: \_\_\_\_\_  
Litigation: \_\_\_\_\_  
Full text: \_\_\_\_\_  
Patent Family: \_\_\_\_\_  
Other: \_\_\_\_\_

VENDOR/COST (where applic.)

STN: \_\_\_\_\_  
DIALOG: \_\_\_\_\_  
Questel/Orbit: \_\_\_\_\_  
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GenCore version 5.1.4 p5\_4578  
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OM protein - protein search, using sw model

Run on: May 6, 2003, 14:55:59 ; Search time 29 Seconds  
(without alignments)  
255.783 Million cell updates/sec

Title: US-09-868-399-1

Perfect score: 198  
Sequence: 1 KIQNFRVYRDSRDLWKGPALKLMKGEGAVVIQDN 36

Scoring table: BLOSUM62  
Gapop 10.0, Gapext 0.5

Searched: 671580 seqs, 206047115 residues

Total number of hits satisfying chosen parameters: 124338

Minimum DB seq length: 0  
Maximum DB seq length: 100

Post-processing: Minimum Match 0%

Maximum Match 100%  
Listing first 45 summaries

Database :

SPTREMBL\_21:\*

- 1: sp\_archaea:\*
- 2: sp\_bacteria:\*
- 3: sp\_fungi:\*
- 4: sp\_human:\*
- 5: sp\_invertebrate:\*
- 6: sp\_mammal:\*
- 7: sp\_mhc:\*
- 8: sp\_organelle:\*
- 9: sp\_phage:\*
- 10: sp\_plant:\*
- 11: sp\_rodent:\*
- 12: sp\_virus:\*
- 13: sp\_vertebrate:\*
- 14: sp\_unclassified:\*
- 15: sp\_virus:\*
- 16: sp\_bacteriap:\*
- 17: sp\_archaeap:\*

Pred. No. is the number of results predicted by chance to have a score greater than or equal to the score of the result being printed, and is derived by analysis of the total score distribution.

## SUMMARIES

Result No.	Score	Query Match	Length	ID	Description
1	147	74.2	90	15	Q88124 chimpanzee
2	147	74.2	90	15	Q88130 chimpanzee
3	112	56.6	93	15	Q87646 chimpanzee
4	56	28.3	59	15	Q90071 chimpanzee
5	53.5	27.0	70	16	P70950 bacillus su
6	51	25.8	96	6	Q9BE76 macaca fasc
7	50.5	25.5	91	16	Q97KS3 clostridium
8	50	25.3	44	15	Q71895 human immun
9	50	25.3	44	15	Q71902 human immun
10	50	25.3	44	15	Q71909 human immun
11	50	25.3	44	15	Q71916 human immun
12	50	25.3	44	15	Q71923 human immun
13	50	25.3	44	15	Q71929 human immun
14	50	25.3	44	15	Q71936 human immun
15	50	25.3	44	15	Q71942 human immun
16	50	25.3	44	15	Q71874 human immun

17	50	25.3	44	15	Q71879 human immun
18	50	25.3	44	15	Q71885 human immun
19	50	25.3	44	15	Q71889 human immun
20	50	25.3	44	15	Q71967 human immun
21	50	25.3	44	15	Q71972 human immun
22	50	25.3	44	15	Q71978 human immun
23	50	25.3	44	15	Q71986 human immun
24	50	25.3	44	15	Q71997 human immun
25	50	25.3	44	15	Q72004 human immun
26	50	25.3	44	15	Q72012 human immun
27	50	25.3	57	5	Q9TX09 geodia cydo
28	49	24.7	77	12	Q8V2L8 camelpox vi
29	48.5	24.5	95	16	Q9X0G0 neisseria m
30	45	22.7	44	15	Q72008 human immun
31	45	22.7	44	15	Q72018 human immun
32	45	22.7	65	15	Q04098 simian t-ce
33	45	22.7	82	16	Q92XC8 rhizobium m
34	44	22.2	44	15	Q71991 human immun
35	44	22.2	80	4	O15521 human sapien
36	43.5	22.0	47	4	Q96CJ4 homocid4
37	43	21.7	52	4	Q9UHS7 homocid4
38	43	21.7	62	4	Q96FS0 homocid4
39	43	21.7	77	13	Q90XF4 coturnix co
40	42.5	21.5	48	11	Q925T2 mus musculu
41	42	21.2	60	16	Q98CP2 rhizobium l
42	42	21.2	76	16	Q8SC0 anabaena sp
43	41.5	21.0	96	16	Q91C5 pseudomonas
44	41	20.7	94	16	Q31639 bacillus su
45	40.5	20.5	38	4	Q9BXG9 homocid4

## ALIGNMENTS

## RESULT 1

Q88124 PRELIMINARY; PRT; 90 AA.

AC Q88124:

DT 01-NOV-1996 (TREMBLrel. 01, Created)

DT 01-NOV-1996 (TREMBLrel. 01, Last sequence update)

DT 01-MAR-2002 (TREMBLrel. 20, Last annotation update)

DE Pol polypeptide (Fragment).

GN GAG-POL OR POL.

OS Chimpanzee immunodeficiency virus (SIV/cpz) (CIV).

CC Viruses; Retroviral viruses; Retroviridae; Lentiviruses.

CC NCBI TaxID=11723;

CC [1]

RP SEQUENCE FROM N.A.

RP MEDLINE=90272009; PubMed=1971917;

RA Dehurs S., Embretson J.E., Anderson D.C., Mullins J.I., Pultz P.N.;

RT "Sequence analysis and acute pathogenicity of molecularly cloned SIV."

RL Nature 345:636-640(1990).

CC -1- PTM: SPECIFIC ENZYMOLOGIC CLEAVAGES IN VIVO YIELD MATURE PROTEINS (BY SIMILARITY).

CC EMBL: I03296; AAA4761.1; -

DR InterPro: IPR001037; Integrase\_C.

DR Pfam: PF00552; Integrase; 1.

KW Endonuclease; Hydrolase; Nucleotidyltransferase; Polypeptide;

FT RNA-directed DNA polymerase.

FT NON TER

FT SEQUENCE 90 AA; 10238 MW; 3FA32AB86D4B57FF CRC64;

Query Match 74.2%; Score 147; DB 15; Length 90;  
Best Local Similarity 70.6%; Pred. No. 1.1e-13;  
Matches 24; Conservative 6; Mismatches 4; Indels 0; Gaps 0;

Cy 1 KIQNFRVYRDSRDLWKGPALKLMKGEGAVVIQ 34  
Db 16 KIQNFRVYRDSRDLWKGPALKLMKGEGAVILK 49

## RESULT 2



RA Boursier R., Boursier L., Brans A., Braun M., Brignell S.C., Bron S.,  
 RA Brouillet S., Bruschi C.V., Caldwell B., Capuano V., Carter N.M.,  
 RA Choi S.K., Codani J.J., Connerion I.F., Cummings N.J., Daniel R.A.,  
 RA Denizot F., Devine K.M., Dusterhoft A., Ehrlich S.D., Emerson P.T.,  
 RA Entlian K.D., Errington J., Fabret C., Ferrari E., Foulger D.,  
 RA Fritz C., Fujita M., Fujita Y., Fuma S., Galizzi A., Galleron N.,  
 RA Gilm S.Y., Glaser P., Goffeau A., Golightly E.J., Grandi G.,  
 RA Giuseppe G., Guy B.J., Haga K., Halech J., Harwood C.R., Henaut A.,  
 RA Hilbert H., Holmappel S., Hosono S., Hullo M.F., Itaya M., Jones L.,  
 RA Joris B., Karamata D., Kanahara Y., Klaer-Blancheard M., Klein C.,  
 RA Kobayashi Y., Koetter P., Koningsstein G., Krogh S., Kumano M.,  
 RA Kurita K., Lapius A., Lardinois S., Lauber J., Lazarevic V.,  
 RA Lee S.M., Levine A., Liu H., Masuda S., Mauei C., Medigic C.,  
 RA Medina N., Meliado R.P., Mizuno M., Moesti D., Nakai S., Pack M.,  
 RA Noone D., O'Reilly M., Ogawa K., Ogiwara A., Oudega B., Park S.H.,  
 RA Paro V., Pohl T.M., Portelle D., Portollik S., Prescott A.M.,  
 RA Pressman E., Pujic P., Putnelle B., Rapoport G., Ray M., Reynolds S.,  
 RA Rieger M., Rivolta C., Roche B., Roche B., Rose W., Sadate Y.,  
 RA Sato T., Scanlan E., Schleich S., Schroeter R., Scoffone F.,  
 RA Sekiguchi J., Sekowska A., Seror S.J., Seror P., Shin B.S., Soldo B.,  
 RA Sorokuch M., Tanakoshi A., Tanaka T., Terpetra P., Tognoni K.,  
 RA Toso V., Uchiyama S., Vandendol M., Vannier F., Vassartot A.,  
 RA Viart A., Wambuit R., Wedler E., Wedler H., Weitzengesser T.,  
 RA Winters P., Wipat A., Yamamoto H., Yamane K., Yasumoto K., Yata K.,  
 RA Yoshida K., Yoshikawa H.F., Zumbstein E., Yoshikawa H., Danchin A.,  
 RT "The complete genome sequence of the gram-positive bacterium *Bacillus*  
 RT subtilis";  
 RL Nature 380:249-256(1997).  
 DR EMBL: Z79580; CAB01839.1; -;  
 DR EMBL: Y09476; CAI70635.1; -;  
 DR EMBL: Z99109; CAB12957.1; -;  
 KM Hypothetical protein; Complete proteome.  
 SQ SEQUENCE 70 AA; 7757 MW; A3490E71E2CDD66 CRC64;

Query Match 27.0%; Score 53.5; DB 16; Length 70;  
 Best Local Similarity 42.9%; Pred. No. 2.8;  
 Matches 12; Conservative 5; Mismatches 10; Indels 1; Gaps 1;

Qy 6 RYVYRDSRPLMKGPALKMKGECAVVI 33  
 Db 44 RLTYR-RRPCKRTGKTIWKNEDAVWV 70

RESULT 6  
 Q9BE76 PRELIMINARY; PRT; 96 AA.  
 AC Q9BE76; 01-JUN-2001 (Tremblrel. 17, Created)  
 DT 01-JUN-2001 (Tremblrel. 17, Last sequence update)  
 DT 01-JUN-2001 (Tremblrel. 17, Last annotation update)  
 DE Hypothetical 10.7 kDa protein.  
 OS Macaca fascicularis (Crab eating macaque) (Cynomolgus monkey).  
 OC Eukaryota; Metazoa; Chordata; Craniata; Vertebrata; Euteleostomi;  
 OC Mammalia; Eutheria; Primates; Catarrhini; Cercopitheidae;  
 OC Cercopitheidae; Macaca.  
 OX NCBI\_TaxID=9541;  
 RN [1]  
 RP SEQUENCE FROM N.A.  
 RC TISSUE=BRAIN PARIETAL LOBE;  
 RA Oada N., Hida M., Kusuda J., Tanuma R., Iseki K., Hirai M., Terao K.,  
 RA Suzuki Y., Sugano S., Hashimoto K.;  
 RT "Isolation of full-length cDNA clones from macaque brain cDNA  
 RT libraries";  
 RL Submitted (MAR-2001) to the EMBL/GenBank/DBJ databases.  
 DR EMBL: AB056770; BAB39318.1; -;  
 KM Hypothetical protein.  
 SQ SEQUENCE 96 AA; 10672 MW; 675945A1C287DA30 CRC64;

Query Match 25.8%; Score 51; DB 6; Length 96;  
 Best Local Similarity 41.7%; Pred. No. 9;  
 Matches 10; Conservative 3; Mismatches 11; Indels 0; Gaps 0;

Qy 10 RDSRDLPLMKGPALKMKGECAVVI 33  
 Db 49 RRSSTPHWQSAHPLMSPGLVLT 72

RESULT 7  
 Q97KS3 PRELIMINARY; PRT; 91 AA.  
 AC Q97KS3; 01-OCT-2001 (Tremblrel. 18, Created)  
 DT 01-OCT-2001 (Tremblrel. 18, Last sequence update)  
 DT 01-JUN-2002 (Tremblrel. 21, Last annotation update)  
 DE Barstar-like protein ribonuclease (barnase) inhibitor.  
 GN CAC0844.  
 OS Clostridium acetobutylicum.  
 OC Bacteria; Firmicutes; Bacillus/Clostridium group; Clostridia;  
 OC Clostridiales; Clostridiaceae; Clostridium.  
 OX NCBI\_TaxID=1488;  
 RN [1]  
 RP SEQUENCE FROM N.A.  
 RC STRAIN=ATCC 824 / DSM 792 / VKM B-1787;  
 RX MEDLINE=21359325; PubMed=11466286;  
 RA Noelling J., Breton G., Omelchenko M.V., Makarova K.S., Zeng Q.,  
 RA Gibson R., Lee H.M., Dubois J., Qiu D., Hiti J., Wolf Y.I.,  
 RA Tatusov R.L., Sabathe F., Doucette-Stamm L., Soucaille P., Daly M.J.,  
 RA Bennett G.N., Koonin E.V., Smith D.R.;  
 RT "Genome sequence and comparative analysis of the solvent-producing  
 RT bacterium *Clostridium acetobutylicum*";  
 RL J. Bacteriol. 183:4823-4838(2001).  
 DR EMBL: AE007600; AAK78820.1; -;  
 DR InterPro: IPR000468; Barstar.  
 DR Pfam: PF01337; Barstar; 1.  
 DR Prodom: PD029050; Barstar; 1.  
 KM Complete proteome.  
 SQ SEQUENCE 91 AA; 10821 MW; AD4B022BC9FCCBBE CRC64;

Query Match 25.5%; Score 50.5; DB 16; Length 91;  
 Best Local Similarity 37.9%; Pred. No. 10;  
 Matches 11; Conservative 3; Mismatches 8; Indels 7; Gaps 1;

Qy 5 FRVYRDSRPLM-----KGPALKMK 26  
 Db 27 FPYYGKNLDMCLTGTETPLKIWK 55

RESULT 8  
 Q71895 PRELIMINARY; PRT; 44 AA.  
 AC Q71895; 01-NOV-1996 (Tremblrel. 01, Created)  
 DT 01-NOV-1996 (Tremblrel. 01, Last sequence update)  
 DT 01-JUN-2002 (Tremblrel. 21, Last annotation update)  
 DE Pol protein (Fragment).  
 GN POL.  
 OS Human immunodeficiency virus type 1.  
 OC Viruses; Retroid viruses; Retroviridae; Lentivirus.  
 OX NCBI\_TaxID=11676;  
 RN [1]  
 RP SEQUENCE FROM N.A.  
 RC STRAIN=CNTRL 1;  
 RX MEDLINE=95287475; PubMed=7769682;  
 RA Michael N.L., Chang G., d'Arcy L.A., Ehrenberg P.K., Mariani R.,  
 RA Busch M.P., Bix D.L., Schwartz D.H.;  
 RT "Defective accessory genes in a human immunodeficiency virus type 1-  
 RT infected long-term survivor lacking recoverable virus";  
 RL J. Virol. 69:4228-4236(1995).  
 DR EMBL: U24447; AAA79545.1; -;  
 DR InterPro: IPR001037; Integrase\_C.  
 DR Pfam: PF00552; Integrase; 1.  
 FT NON\_TER 1 1  
 SQ SEQUENCE 44 AA; 4875 MW; 01901EA27AFA3CDE CRC64;

Query Match 25.3%; Score 50; DB 15; Length 44;

Best Local Similarity 100.0%; Pred. No. 5.3;  
Matches 10; Conservative 0; Mismatches 0;

QY 27 GEGAVI0DN 36  
| | | | | | | | | |  
DB 1 GEGAVI0DN 10

## RESULT 9

ID Q71902 PRELIMINARY; PRT; 44 AA.  
AC Q71902;  
DT 01-NOV-1996 (TREMBlrel. 01, Created)  
DT 01-NOV-1996 (TREMBlrel. 01, Last sequence update)  
DE 01-JUN-2002 (TREMBlrel. 21, Last annotation update)  
GN POL.  
OS Human immunodeficiency virus type 1.  
OC Viruses; Retrovirdae; Retroviridae; Lentivirus.  
OX NCBI\_TaxID=11676;  
RN [1]  
RP SEQUENCE FROM N.A.  
RC STRAIN=CNTRL.1;  
RX MEDLINE=95287475; PubMed=7769682;  
RA Michael N.L., Chang G., d'Arcy L.A., Ehrenberg P.K., Mariani R.,  
RT "Defective accessory genes in a human immunodeficiency virus type 1-  
RL J. Virol. 69:4228-4236(1995).  
DR EMBL; U24449; AAA79552.1; -  
DR InterPro: IPR001037; Integrase\_C.  
DR Pfam: PF00552; Integrase; 1.  
FT NON TER  
SQ SEQUENCE 44 AA; 4875 MW; 01901EA27AFA3CDE CRC64;

Query Match 25.3%; Score 50; DB 15; Length 44;  
Best Local Similarity 100.0%; Pred. No. 5.3;  
Matches 10; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

QY 27 GEGAVI0DN 36  
| | | | | | | | | |  
DB 1 GEGAVI0DN 10

## RESULT 10

ID Q71909 PRELIMINARY; PRT; 44 AA.  
AC Q71909;  
DT 01-NOV-1996 (TREMBlrel. 01, Created)  
DT 01-NOV-1996 (TREMBlrel. 01, Last sequence update)  
DE 01-JUN-2002 (TREMBlrel. 21, Last annotation update)  
GN POL.  
OS Human immunodeficiency virus type 1.  
OC Viruses; Retrovirdae; Retroviridae; Lentivirus.  
OX NCBI\_TaxID=11676;  
RN [1]  
RP SEQUENCE FROM N.A.  
RC STRAIN=CNTRL.1;  
RX MEDLINE=95287475; PubMed=7769682;  
RA Michael N.L., Chang G., d'Arcy L.A., Ehrenberg P.K., Mariani R.,  
RT "Defective accessory genes in a human immunodeficiency virus type 1-  
RL J. Virol. 69:4228-4236(1995).  
DR EMBL; U24449; AAA79552.1; -  
DR InterPro: IPR001037; Integrase\_C.  
DR Pfam: PF00552; Integrase; 1.  
FT NON TER  
SQ SEQUENCE 44 AA; 4875 MW; 01901EA27AFA3CDE CRC64;

Query Match 25.3%; Score 50; DB 15; Length 44;  
Best Local Similarity 100.0%; Pred. No. 5.3;

Matches 10; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

QY 27 GEGAVI0DN 36  
| | | | | | | | | |  
DB 1 GEGAVI0DN 10

## RESULT 11

ID Q71916 PRELIMINARY; PRT; 44 AA.  
AC Q71916;  
DT 01-NOV-1996 (TREMBlrel. 01, Created)  
DT 01-NOV-1996 (TREMBlrel. 01, Last sequence update)  
DE 01-JUN-2002 (TREMBlrel. 21, Last annotation update)  
GN POL.  
OS Human immunodeficiency virus type 1.  
OC Viruses; Retrovirdae; Retroviridae; Lentivirus.  
OX NCBI\_TaxID=11676;  
RN [1]  
RP SEQUENCE FROM N.A.  
RC STRAIN=CNTRL.1;  
RX MEDLINE=95287475; PubMed=7769682;  
RA Michael N.L., Chang G., d'Arcy L.A., Ehrenberg P.K., Mariani R.,  
RT "Defective accessory genes in a human immunodeficiency virus type 1-  
RL J. Virol. 69:4228-4236(1995).  
DR EMBL; U24451; AAA79573.1; -  
DR InterPro: IPR001037; Integrase\_C.  
DR Pfam: PF00552; Integrase; 1.  
FT NON TER  
SQ SEQUENCE 44 AA; 4875 MW; 01901EA27AFA3CDE CRC64;

Query Match 25.3%; Score 50; DB 15; Length 44;  
Best Local Similarity 100.0%; Pred. No. 5.3;  
Matches 10; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

QY 27 GEGAVI0DN 36  
| | | | | | | | | |  
DB 1 GEGAVI0DN 10

## RESULT 12

ID Q71923 PRELIMINARY; PRT; 44 AA.  
AC Q71923;  
DT 01-NOV-1996 (TREMBlrel. 01, Created)  
DT 01-NOV-1996 (TREMBlrel. 01, Last sequence update)  
DE 01-JUN-2002 (TREMBlrel. 21, Last annotation update)  
GN POL.  
OS Human immunodeficiency virus type 1.  
OC Viruses; Retrovirdae; Retroviridae; Lentivirus.  
OX NCBI\_TaxID=11676;  
RN [1]  
RP SEQUENCE FROM N.A.  
RC STRAIN=CNTRL.1;  
RX MEDLINE=95287475; PubMed=7769682;  
RA Michael N.L., Chang G., d'Arcy L.A., Ehrenberg P.K., Mariani R.,  
RT "Defective accessory genes in a human immunodeficiency virus type 1-  
RL J. Virol. 69:4228-4236(1995).  
DR EMBL; U24451; AAA79573.1; -  
DR InterPro: IPR001037; Integrase\_C.  
DR Pfam: PF00552; Integrase; 1.  
FT NON TER  
SQ SEQUENCE 44 AA; 4875 MW; 01901EA27AFA3CDE CRC64;

Query Match 25.3%; Score 50; DB 15; Length 44;  
Best Local Similarity 100.0%; Pred. No. 5.3;  
Matches 10; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

Oy 27 GEGAVI0DN 36  
|||||  
Db 1 GEGAVI0DN 10

## RESULT 13

Q71929 PRELIMINARY; PRT; 44 AA.  
ID 071929  
AC 071929  
DT 01-NOV-1996 (TrEMBLrel. 01, Created)  
DT 01-NOV-1996 (TrEMBLrel. 01, Last sequence update)  
DT 01-JUN-2002 (TrEMBLrel. 21, Last annotation update)  
DE Pol protein (Fragment).  
GN POL.  
OS Human immunodeficiency virus type 1.  
OC Viruses; Retroid viruses; Retroviridae; Lentivirus.  
OX NCBI\_TaxID=11676;  
RN [1]  
RP SEQUENCE FROM N.A.  
RC STRAIN=CNTRL.1;  
RX MEDLINE=95287475; PubMed=7769682;  
RA Michael N.L., Chang G., d'Arcy L.A., Ehrenberg P.K., Mariani R.,  
RA Busch M.P., Birx D.L., Schwartz D.H.;  
RT "Defective accessory genes in a human immunodeficiency virus type 1-  
RT infected long-term survivor lacking recoverable virus.";  
RL J. Virol. 69:4228-4236(1995).  
DR EMBL: U24452; AAA79579.1;  
DR InterPro: IPR001037; Integrase\_C.  
DR Pfam: PF00552; Integrase; 1.  
FT NON TER 1  
SQ SEQUENCE 44 AA; 4875 MW; 01901EA27AFA3CDE CRC64;

Query Match 25.3%; Score 50; DB 15; Length 44;  
Best Local Similarity 100.0%; Pred. No. 5.3;  
Matches 10; Conservative 0; Mismatches 0; Indels 0; Gaps 0;  
Oy 27 GEGAVI0DN 36  
|||||  
Db 1 GEGAVI0DN 10

## RESULT 14

Q71936 PRELIMINARY; PRT; 44 AA.  
ID 071936  
AC 071936  
DT 01-NOV-1996 (TrEMBLrel. 01, Created)  
DT 01-NOV-1996 (TrEMBLrel. 01, Last sequence update)  
DT 01-JUN-2002 (TrEMBLrel. 21, Last annotation update)  
DE Pol protein (Fragment).  
GN POL.  
OS Human immunodeficiency virus type 1.  
OC Viruses; Retroid viruses; Retroviridae; Lentivirus.  
OX NCBI\_TaxID=11676;  
RN [1]  
RP SEQUENCE FROM N.A.  
RC STRAIN=CNTRL.2;  
RX MEDLINE=95287475; PubMed=7769682;  
RA Michael N.L., Chang G., d'Arcy L.A., Ehrenberg P.K., Mariani R.,  
RA Busch M.P., Birx D.L., Schwartz D.H.;  
RT "Defective accessory genes in a human immunodeficiency virus type 1-  
RT infected long-term survivor lacking recoverable virus.";  
RL J. Virol. 69:4228-4236(1995).  
DR EMBL: U24453; AAA79586.1;  
DR InterPro: IPR001037; Integrase\_C.  
DR Pfam: PF00552; Integrase; 1.  
FT NON TER 1  
SQ SEQUENCE 44 AA; 4690 MW; F0F71C7266DF3CC6 CRC64;

Query Match 25.3%; Score 50; DB 15; Length 44;  
Best Local Similarity 100.0%; Pred. No. 5.3;  
Matches 10; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

Oy 27 GEGAVI0DN 36  
|||||  
Db 1 GEGAVI0DN 10

## RESULT 15

Q71942 PRELIMINARY; PRT; 44 AA.  
ID 071942  
AC 071942  
DT 01-NOV-1996 (TrEMBLrel. 01, Created)  
DT 01-NOV-1996 (TrEMBLrel. 01, Last sequence update)  
DT 01-JUN-2002 (TrEMBLrel. 21, Last annotation update)  
DE Pol protein (Fragment).  
GN POL.  
OS Human immunodeficiency virus type 1.  
OC Viruses; Retroid viruses; Retroviridae; Lentivirus.  
OX NCBI\_TaxID=11676;  
RN [1]  
RP SEQUENCE FROM N.A.  
RC STRAIN=CNTRL.2;  
RX MEDLINE=95287475; PubMed=7769682;  
RA Michael N.L., Chang G., d'Arcy L.A., Ehrenberg P.K., Mariani R.,  
RA Busch M.P., Birx D.L., Schwartz D.H.;  
RT "Defective accessory genes in a human immunodeficiency virus type 1-  
RT infected long-term survivor lacking recoverable virus.";  
RL J. Virol. 69:4228-4236(1995).  
DR EMBL: U24454; AAA79592.1;  
DR InterPro: IPR001037; Integrase\_C.  
DR Pfam: PF00552; Integrase; 1.  
FT NON TER 1  
SQ SEQUENCE 44 AA; 4762 MW; F15C041266DF3CC6 CRC64;

Query Match 25.3%; Score 50; DB 15; Length 44;  
Best Local Similarity 100.0%; Pred. No. 5.3;  
Matches 10; Conservative 0; Mismatches 0; Indels 0; Gaps 0;  
Oy 27 GEGAVI0DN 36  
|||||  
Db 1 GEGAVI0DN 10

Search completed: May 6, 2003, 14:58:17  
Job time : 31 secs





GenCore version 5.1.4 p5 4578  
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OM protein - protein search, using sw model

Run on: May 6, 2003, 14:55:39 / Search time 11 Seconds

(without alignments)  
135.741 Million cell updates/sec

Title: US-09-868-399-1

Perfect score: 198  
Sequence: 1 KIQNFRVRYSDRDLWKGPALKLWKGEAVIQDN 36

Scoring table: BIOSUM62  
Gapop 10.0, Gapext 0.5

Searched: 112892 seqs, 41476328 residues

Total number of hits satisfying chosen parameters: 12886

Minimum DB seq length: 0

Maximum DB seq length: 100

Post-processing: Minimum Match 0%

Maximum Match 100%

Listing first 45 summaries

Database: SwissProt\_40\*

Pred. No. is the number of results predicted by chance to have a score greater than or equal to the score of the result being printed, and is derived by analysis of the total score distribution.

## SUMMARIES

- Result No.	Score	Query Match	Length	ID	Description
1	38.5	19.4	96	1	P71836 synechocyst
2	38	19.3	83	1	VG03_BPM2
3	37.5	18.9	73	1	GBG8_HUMAN
4	36.5	18.4	97	1	IFIA_ARCFU
5	36	18.2	36	1	VG50_HAELN
6	36	18.2	73	1	GBG1_BOVIN
7	36	18.2	87	1	R35A_PYRAB
8	36	18.2	87	1	R35A_PYRFU
9	36	18.2	87	1	R35A_PYRMO
10	36	18.2	89	1	VYAP_VACCC
11	35.5	17.9	89	1	BARS_BACAM
12	35.5	17.9	91	1	SR19_METTH
13	35	17.7	76	1	A4_MACMU
14	35	17.7	83	1	VG7_SPLVR
15	35	17.7	85	1	VG73_BPM5
16	35	17.7	87	1	A4_MACFA
17	34.5	17.4	73	1	VB76_HAELN
18	34.5	17.4	97	1	Y117_NPVOF
19	34	17.2	56	1	AMCI_APIPE
20	34	17.2	65	1	LHA2_ECTHL
21	34	17.2	66	1	LYED_VACCC
22	34	17.2	88	1	V4OL_RHTSN
23	34	17.2	93	1	PFTB_PSEAE
24	34	17.2	97	1	YBGE_ECOLI
25	33.5	16.9	79	1	RS16_BUCAI
26	33.5	16.9	82	1	RS16_YERPE
27	33.5	16.9	92	1	RS19_CAUCR
28	33.5	16.9	93	1	RT19_MARPO
29	33.5	16.9	94	1	ES65_MYCTU
30	33.5	16.9	94	1	ES66_MYCTU
31	33	16.7	65	1	PSAI_ANAVA
32	33	16.7	71	1	HXC5_NORVI
33	33	16.7	75	1	MT1B_VICFA

34	33	16.7	77	1	VG43_BPM5	Q05255 mycobacteri
35	33	16.7	83	1	Y67_BPT3	P20330 bacterioph
36	33	16.7	92	1	RS19_RICCN	Q92930 rickettsia
37	33	16.7	92	1	RS19_RICPR	Q92939 rickettsia
38	32.5	16.4	36	1	NUCM_SOLTU	Q02654 solanum tub
39	32.5	16.4	92	1	RS19_AGR75	Q04252 agrobacteri
40	32.5	16.4	94	1	IFAZ_HAELN1	Q04257 halobacteri
41	32.5	16.4	98	1	S10Z_HUMAN	Q04258 homo sapien
42	32	16.2	17	1	PH4_PBRAM	P82697 periplaneta
43	32	16.2	37	1	ES2E_RANES	P40846 rana escul
44	32	16.2	52	1	A1T3_HORSE	P38030 equus cabal
45	32	16.2	.58	1	68MP_MOUSE	P56379 mus muscul

## ALIGNMENTS

RESULT 1	YV02_SYNY3	STANDARD;	PRT;	96 AA.
ID	P71836			
AC	15-JUL-1998 (Rel. 36, Created)			
DT	15-JUL-1998 (Rel. 36, Last sequence update)			
DE	Hypothetical protein ser3402 precursor.			
GN	SSR3402.			
OS	Synechocystis sp. (strain PCC 6803).			
OC	Bacteria; Cyanobacteria; Chroococcales; Synechocystis.			
OX	NCBI_Taxid=1148;			
ON	[1]			
RP	SEQUENCE FROM N.A.			
RX	MEDLINE=97061201; PubMed=8905231;			
RA	Kaneko T., Sato S., Kotani H., Tanaka A., Asamizu E., Nakamura Y.,			
RA	Myajima N., Hirosewa M., Sugita M., Sasamoto S., Kimura T.,			
RA	Hosouchi T., Matsuno A., Muraki A., Nakazaki N., Natsu K., Okumura S.,			
RA	Shimpo S., Takeuchi C., Wada T., Watanabe A., Yamada M., Yasuda M.,			
RA	Tabata S.;			
RT	"Sequence analysis of the genome of the unicellular cyanobacterium			
RT	Synechocystis sp. strain PCC6803. II. Sequence determination of the			
RT	entire genome and assignment of potential protein-coding regions."			
RL	DNA Res. 3:109-136(1996).			
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CC	or send an email to <a href="mailto:license@isb-sib.ch">license@isb-sib.ch</a> ).			
CC	-----			
DR	EMBL, D90910, BAI17893.1, -;			
KW	Hypothetical protein; Signal; Complete proteome.			
FT	SIGNAL			
FT	CHAIN 20 96			
FT	CHAIN 1 19			
SQ	SEQUENCE 96 AA; 10673 MW; 7417586D70068C3E CIG64;			
Query Match	Score 38.5; DB 1; Length 96;			
Best Local Similarity	56.2%; Pred. No. 1.2e+02;			
Matches	9; Conservative 2; Mismatches 4; Indels 1; Gaps 1;			
Db	10 RDSRPL-WKGPALKL 24			
	71 KPSKPLWAGPALKL 86			
RESULT 2	VG03_BPM2	STANDARD;	PRT;	83 AA.
ID	OG4159;			
AC	15-DEC-1998 (Rel. 37, Created)			
DT	15-DEC-1998 (Rel. 37, Last sequence update)			
DE	15-DEC-1998 (Rel. 37, Last annotation update)			
DE	Gene 3 protein (Gp3).			

```

GN OS Mycobacteriophage D29.
OC Viruses: dsDNA viruses, no RNA stage; Caudovirales; Siphoviridae;
OC NCBI_TaxID=28369.
OX NCBI_TaxID=28369.
RN [1]
RP SEQUENCE FROM N.A.
RX MEDLINE=98300335; PubMed=9636706.
RA Ford M.E., Sarkis G.J., Belanger A.B., Hendrix R.W., Hatfull G.F.;
RT "Genome structure of mycobacteriophage D29: implications for phage
RL J. Mol. Biol. 279:143-164(1998).
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CC or send an email to license@isb-sib.ch).
DR EMBL AF022214; AAC18446.1;
SQ SEQUENCE 83 AA; 8962 MW; 2B3D070B4C6F4E09 CRC64;

Query Match 19.2%; Score 38; DB 1; Length 83;
Best Local Similarity 36.8%; Pred. No. 1.2e+02;
Matches 7; Conservative 4; Mismatches 8; Indels 0; Gaps 0;

OY 17 WKGPRLMKRGAGAVI00 35
Db 42 WEGLEILEYSGDGYEVSD 60

RESULT 3
GGBB HUMAN
AC GGBB HUMAN STANDARD; PRT; 73 AA.
AC P50152:
DT 01-OCT-1996 (Rel. 34, Created)
DT 01-OCT-1996 (Rel. 34, Last sequence update)
DT 15-JUN-2002 (Rel. 41, Last annotation update)
GN Gamma nucleotide-binding protein G(1)/G(S)/G(O) gamma-11 subunit.
DS GNG11 OR GNG11.
OS Homo sapiens (Human).
OS Mus musculus (Mouse), and
OS Rattus norvegicus (Rat).
OC Eukaryota; Metazoa; Chordata; Craniata; Vertebrata; Euteleostomi;
OC Mammalia; Eutheria; Primates; Catarrhini; Homiidae; Homo.
OX NCBI_TaxID=9606, 10090, 10116,
RN [1]
RP SEQUENCE FROM N.A., AND ISOPRENOID.
RC SPECIES=Human; TISSUE=Testis;
RX MEDLINE=95394940; PubMed=765396;
RA Ray K., Kunsch C., Bonner L.M., Robishaw J.D.;
RT "Isolation of cDNA clones encoding eight different human G protein
RT gamma subunits, including three novel forms designated the gamma 4,
RL J. Biol. Chem. 270:21765-21771(1995).
RN [2]
RP SEQUENCE FROM N.A.
RC SPECIES=Human;
RA Maggi L.;
RL Submitted (MAY-1997) to the EMBL/GenBank/DBJ databases.
RN [3]
RP SEQUENCE FROM N.A.
RC SPECIES=Human; TISSUE=Lung;
RA Straube R.;
RL Submitted (JUN-2001) to the EMBL/GenBank/DBJ databases.
RN [4]
RP SEQUENCE FROM N.A.
RC SPECIES=Mouse; STRAIN=C57BL/6J; TISSUE=Kidney, and Tongue;
RX MEDLINE=21085660; PubMed=11217851;
RA Kawai J., Shingawa A., Shibata K., Yoshino M., Itoh M., Ishi Y.,
RA Atakawa T., Hara A., Fukunishi Y., Kono H., Adachi J., Fukuda S.,

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RA Alizawa K., Izawa M., Nishi K., Kiyosawa H., Kondo S., Yamanaka I.,
RA Saito T., Okazaki Y., Gotohori T., Bono H., Kasukawa T., Saito R.,
RA Kadota K., Matsuda H.A., Ashburner M., Batilov S., Casavant T.,
RA Fleischmann W., Gaasterland T., Gissi C., King B., Kochava H.,
RA Kuehl P., Lewis S., Matsuo Y., Nikaido I., Pesole G., Quackenbush J.,
RA Schirml L.M., Staudl F., Suzuki R., Tomita M., Wagner L., Washio T.,
RA Sakai K., Okido T., Furuno M., Aono H., Baldarelli R., Barsh G.,
RA Blake J., Boffelli D., Bojunga N., Carninci P., de Bonaldo M.F.,
RA Brownstein M.J., Bull C., Fletcher C., Fujita M., Gariboldi M.,
RA Gustincich S., Hill D., Hofmann M., Hume D.A., Kamiya M., Lee N.H.,
RA Lyons P., Marchionni L., Mashima J., Mazzarelli J., Mombaerts P.,
RA Nordone P., Ring B., Ringwald M., Rodriguez I., Sakamoto N.,
RA Sasaki H., Sato K., Schoenbach C., Seya T., Shibata Y., Storch K.-F.,
RA Suzuki K., Toyooka K., Wang K.H., Weitz C., Whitaker C., Wilming L.,
RA Wynshaw-Boris A., Yoshida K., Hasegawa Y., Kawai H., Kohlsuki S.,
RA Hayashizaki Y.;
RT "Functional annotation of a full-length mouse cDNA collection.";
RL Nature 409:685-690(2001).
RN [5]
RP SEQUENCE FROM N.A.
RC SPECIES=Rat; STRAIN=Sprague-Dawley;
RA Costain W.J., Mishra R.K.;
RT "Identification and cloning of rat G protein gamma 11 subunit.";
RL Submitted (APR-2000) to the EMBL/GenBank/DBJ databases.
CC -1- FUNCTION: GUANINE NUCLEOTIDE-BINDING PROTEIN (G PROTEIN) ARE
CC INVOLVED AS A MODULATOR OR TRANSDUCER IN VARIOUS TRANSMEMBRANE
CC SIGNALING SYSTEMS. THE BETA AND GAMMA CHAINS ARE REQUIRED FOR THE
CC GTPASE ACTIVITY, FOR REPLACEMENT OF GDP BY GTP, AND FOR G PROTEIN-
CC RECEPTOR INTERACTION. INTERACTS WITH BETA-1 AND BETA-3, BUT NOT
CC WITH BETA-2.
CC -1- SUBUNIT: G PROTEINS ARE COMPOSED OF 3 UNITS (ALPHA, BETA & GAMMA).
CC -1- TISSUE SPECIFICITY: ABUNDANTLY EXPRESSED IN ALL TISSUES TESTED
CC EXCEPT FOR BRAIN.
CC -1- SIMILARITY: BELONGS TO THE G PROTEIN GAMMA FAMILY.
CC
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DR EMBL, U31384; AAC50206.1;
DR EMBL, AC002076; AAB54051.1;
DR EMBL, BC009709; AAB09709.1;
DR EMBL, AK002765; BAB2340.1;
DR EMBL, AK009529; BAB26342.1;
DR EMBL, AF257110; AAF68984.1;
DR HSSP; P02698; 189X.
DR GeneW; HGNC:4403; GNG11.
DR MIM; 604390;
DR InterPro; IPR001770; G-gamma.
DR InterPro; IPR001230; Prenyl_site.
DR Pfam; PF00631; G-gamma; 1.
DR PRINTS; PR00321; GPROTEING.
DR PRODom; PD003763; G-gamma; 1.
DR SMART; SM00224; GGL; 1.
DR PROSITE; PS50058; G_PROTEIN_GAMMA; 1.
DR Transducer; Prenylation; Lipoprotein; Multigene family.
FT LIPID 70
FT LIPID 70
FT PROPEP 71
SQ SEQUENCE 73 AA; 8481 MW; 2B13935E3AEFB9E8 CRC64;

Query Match 18.9%; Score 37.5; DB 1; Length 73;
Best Local Similarity 37.1%; Pred. No. 1.2e+02;
Matches 13; Conservative 7; Mismatches 12; Indels 3; Gaps 2;

OY 1 KIONPRVYRDSRDLPMG--PAKLWKEGAGAVI 33
Db 39 EIKNY-IEBRSGEPLVKGIPEDKNPFKEKSCVI 72

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RESULT 4
ID IF1A_ARCFU STANDARD; PRT; 97 AA.
AC 029481;
DT 15-JUL-1998 (Rel. 36, Created)
DT 16-OCT-2001 (Rel. 40, Last sequence update)
DT 15-JUN-2002 (Rel. 41, Last annotation update)
DE Probable translation initiation factor 1A (aIF-1A).
GN aIF1A OR AF0777.
OS Archaeoglobus fulgidus.
OC Archaea; Euryarchaeota; Archaeoglobi; Archaeoglobales;
OC Archaeoglobaceae; Archaeoglobus.
OX NCBI_TaxID=2234;
RN [1]
RP SEQUENCE FROM N.A.
RC STRAIN=VC-16 / DSM 4304 / ATCC 49558;
RX MEDLINE=98049343; PubMed=9389475;
RA Kleink H.-P., Clayton R.A., Tomb J.-F., White O., Nelson K.E.,
RA Ketchum K.A., Dodson R.J., Gwinn M., Hickley E.K., Peterson J.D.,
RA Richardson D.L., Kerkvliet A.R., Graham D.E., Kyriades N.C.,
RA Fleischmann R.D., Quackenbush J., Lee N.H., Sutton G.G., Gill S.,
RA Kirkness E.F., Dougherty B.A., McKenney K., Adams M.D., Loftus B.,
RA Peterson S., Reich C.I., McNeil L.K., Badger J.H., Glodek A., Zhou L.,
RA Overbeek R., Gocayne J.D., Weidman J.F., McDonald L., Uterback T.,
RA Cotton M.D., Spriggs T., Artlich P., Kaine B.P., Sykes S.M.,
RA Sadow P.W., D'Andrea K.P., Bowman C., Fujii C., Garland S.A.,
RA Mason T.M., Olsen G.J., Fraser C.M., Smith H.O., Woese C.R.,
RA Venter J.C.;
RT "The complete genome sequence of the hyperthermophilic, sulphate-
RT reducing archaeon Archaeoglobus fulgidus.";
RL Nature 390:364-370(1997).
CC -1- FUNCTION: Seems to be required for maximal rate of protein
CC biosynthesis. Enhances ribosome dissociation into subunits and
CC stabilizes the binding of the initiator Met-tRNA(I) to 40 S
CC ribosomal subunits (By similarity).
CC -1- SIMILARITY: BELONGS TO THE EIF-1A FAMILY.
CC -1- SIMILARITY: CONTAINS 1 SL-LIKE DOMAIN.
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CC -----
DR EMBL; AE001051; AAB90469.1; ALT_INIT.
DR TIGR; AF0777; -.
DR InterPro; IPR001253; TIF_EIF-1A.
DR Pfam; PF01176; eIF-1a; 1.
DR ProDom; PD005579; TIF_EIF-1A; 1.
DR TrRfam; TRF00523; eIF-1A; 1.
DR PROSITE; PS01262; IF1A; 1.
DR PROSITE; PS50832; SL-IF1_TYPE; 1.
KM Initiation factor; Protein biosynthesis; Complete proteome.
FT DOMAIN 8 82 SL-LIKE.
SQ SEQUENCE 97 AA; 11385 MW; 12332D5E9CB82DA1 CRC64;

Query Match 18.4%; Score 36.5; DB 1; Length 97;
Best local Similarity 33.3%; Pred. No. 2.2e+02;
Matches 11; Conservative 6; Mismatches 13; Indels 3; Gaps 1;
```

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DT 15-JUN-2002 (Rel. 41, Last annotation update)
DE Hypothetical protein H1650.
GN H1650.
OS Haemophilus influenzae.
OC Bacteria; Proteobacteria; gamma subdivision; Pasteurellaceae;
OC Haemophilus.
OX NCBI_TaxID=727;
RN [1]
RP SEQUENCE FROM N.A.
RC STRAIN=Rd / KW20 / ATCC 51907;
RX MEDLINE=95350630; PubMed=7542800;
RA Fleischmann R.D., Adams M.D., White O., Clayton R.A., Kirkness E.F.,
RA Kerkvliet A.R., Bult C.J., Tomb J.-F., Dougherty B.A., Merrick J.M.,
RA McKenney K., Sutton G., Fitzhugh W., Fields C.A., Gocayne J.D.,
RA Scott J.D., Shirley R., Liu L.-I., Glodek A., Kelley J.M.,
RA Weidman J.F., Phillips C.A., Spriggs T., Hedblom E., Cotton M.D.,
RA Uterback T.R., Hanna M.C., Nguyen D.T., Sauder D.M., Brandon R.C.,
RA Fine L.D., Fritchman J.L., Fuhrmann J.L., Geoghagen N.S.M.,
RA Gnehm C.L., McDonald L.A., Small K.V., Fraser C.M., Smith H.O.,
RA Venter J.C.;
RT "Whole-genome random sequencing and assembly of Haemophilus influenzae
RT Rd.";
RL Science 269:496-512(1995).
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CC -----
DR EMBL; U32838; AAC23303.1; -.
DR TIGR; H1650; -.
RW Hypothetical protein; Complete proteome.
SQ SEQUENCE 36 AA; 4285 MW; 248010DAD7898B33 CRC64;

Query Match 18.2%; Score 36; DB 1; Length 36;
Best local Similarity 42.9%; Pred. No. 90;
Matches 9; Conservative 2; Mismatches 8; Indels 2; Gaps 1;

Db 7 IONHSTYNNRD-WIGYOK 25

RESULT 6
GBG1_BOVIN STANDARD; PRT; 73 AA.
ID GBG1_BOVIN
AC P02698;
DT 21-JUL-1986 (Rel. 01, Created)
DT 21-JUL-1986 (Rel. 01, Last sequence update)
DT 16-OCT-2001 (Rel. 40, Last annotation update)
DE Guanine nucleotide-binding protein G(T) gamma-T1 subunit (Transducin
DE gamma chain).
GN GNGT1.
OS Bos taurus (Bovine).
OC Eukaryota; Metazoa; Chordata; Craniata; Vertebrata; Euteleostomi;
OC Mammalia; Eutheria; Cetartiodactyla; Ruminantia; Pecora; Bovidae;
OC Bovidae; Bovinae; Bos.
OX NCBI_TaxID=9913;
RN [1]
RP SEQUENCE FROM N.A.
RX MEDLINE=85063709; PubMed=6438626;
RA Hurley J.B., Fong H.K.W., Teplow D.B., Dreyer W.J., Simon M.I.;
RT "Isolation and characterization of a cDNA clone for the gamma subunit
RT of bovine retinal transducin.";
RL Proc. Natl. Acad. Sci. U.S.A. 81:6948-6952(1984).
RN [2]
RP SEQUENCE FROM N.A.
RX MEDLINE=85166247; PubMed=2984674;
RA Yatsunami K., Pandya B.V., Orian D.D., Khorana H.G.;
RT "cDNA-derived amino acid sequence of the gamma subunit of GTPase from
```

RT bovine rod outer segments.";  
 RL Proc. Natl. Acad. Sci. U.S.A. 82:1936-1940(1985).  
 RN [3]  
 RP SEQUENCE FROM N.A.  
 RA MEDLINE=93272877; PubMed=8500562;  
 RT Tao L., Pandey S., Simon M.I., Fong H.K.;  
 RT "Structure of the bovine transducin gamma subunit gene and analysis  
 RT of promoter function in transgenic mice.";  
 RN Exp. Eye Res. 56:497-507(1993).  
 [4]  
 RP SEQUENCE OF 1-39 FROM N.A.  
 RA MEDLINE=85046503; PubMed=6149748;  
 RT van Dorp C., Medynski D.C., Sullivan K., Wu A.M., Fung B.K.-K.,  
 RT Bourne H.R.;  
 RT "Partial cDNA sequence of the gamma subunit of transducin.";  
 RL Biochem. Biophys. Res. Commun. 124:250-255(1984).  
 [5]  
 RP SEQUENCE OF 1-69.  
 RA MEDLINE=85076983; PubMed=3917402;  
 RT Ovchinnikov Y.A., Lipkin V.M., Shvavaeva T.M., Bogachuk A.P.,  
 RT Shenyakin V.V.;  
 RT "Complete amino acid sequence of gamma-subunit of the GTP-binding  
 RT protein from cattle retina.";  
 RL FEBS Lett. 179:107-110(1985).  
 [6]  
 RP ISOPRENOID.  
 RA MEDLINE=90348966; PubMed=2385292;  
 RT Fukuda Y., Takao T., Ohguro H., Yoshizawa T., Akino T.,  
 RT Shimomishi Y.;  
 RT "Farnesylated gamma-subunit of photoreceptor G protein indispensable  
 RT for GTP-binding.";  
 RL Nature 346:658-660(1990).  
 [7]  
 RP ISOPRENOID.  
 RA MEDLINE=91236727; PubMed=1903391;  
 RT Sanford J., Codina J., Birnbaumer L.;  
 RT "Gamma-subunits of G proteins, but not their alpha- or beta-subunits,  
 RT are polyisoprenylated. Studies on post-translational modifications  
 RT using in vitro translation with rabbit reticulocyte lysates.";  
 RL J. Biol. Chem. 266:9570-9579(1991).  
 [8]  
 RP ISOPRENOID.  
 RA MEDLINE=91017567; PubMed=2217200;  
 RT Lai R.K., Perez-Sala D., Canada F.J., Rando R.R.;  
 RT "The gamma subunit of transducin is farnesylated.";  
 RL Proc. Natl. Acad. Sci. U.S.A. 87:7673-7677(1990).  
 [9]  
 RP X-RAY CRYSTALLOGRAPHY (2.1 ANGSTROMS) OF BETA-GAMMA DIMER.  
 RA MEDLINE=96149254; PubMed=8552196;  
 RT Sander J., Bohm A., Lambright D.G., Hamm H.E., Sigler P.B.;  
 RT "Crystal structure of a G-protein beta gamma dimer at 2.1-A  
 RT resolution.";  
 RL Nature 379:369-374(1996).  
 [10]  
 RP X-RAY CRYSTALLOGRAPHY (2.8 ANGSTROMS) OF COMPLEX WITH PHOSUDCIN.  
 RA MEDLINE=98416696; PubMed=9739091;  
 RT Loew A., Ho Y.K., Blundell T., Bax B.;  
 RT "Phosducin induces a structural change in transducin beta gamma.";  
 RL Structure 6:1007-1019(1998).  
 [11]  
 RP FUNCTION: GUANINE NUCLEOTIDE-BINDING PROTEINS (G PROTEINS) ARE  
 RP INVOLVED AS A MODULATOR OR TRANSDUCER IN VARIOUS TRANSMEMBRANE  
 RP SIGNALING SYSTEMS. THE BETA AND GAMMA CHAINS ARE REQUIRED FOR THE  
 RP EFFECTOR INTERACTION.  
 [12]  
 RP SUBUNIT: G PROTEINS ARE COMPOSED OF 3 UNITS (ALPHA, BETA & GAMMA).  
 [13]  
 RP TISSUE SPECIFICITY: RETINAL ROD OUTER SEGMENT.  
 [14]  
 RP SIMILARITY: BELONGS TO THE G PROTEIN GAMMA FAMILY.  
 [15]  
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 CC -----  
 DR EMBL: K03255; AAA0794.1; -  
 CC EMBL: K02199; AAA0793.1; -  
 DR EMBL: S62031; AAB26895.1; -  
 CC EMBL: S62029; AAB26895.1; JOINED.  
 DR EMBL: K02436; AAA0788.1; -  
 CC PIR: A03153; RGHOGT  
 DR PDB: 1TRG; 01-APR-97.  
 DR PDB: 1AOR; 16-FEB-99.  
 DR PDB: 1B9X; 23-FEB-99.  
 DR PDB: 1B9Y; 23-FEB-99.  
 DR InterPro: IPR001770; G-gamma.  
 DR InterPro: IPR001230; Prenyl\_site.  
 DR Pfam: PF00631; G-gamma; 1.  
 DR PRINTS: PR00321; GPROTEIN.  
 DR ProDom: PD003783; G-gamma; 1.  
 DR SMART: SM00224; GGL; 1.  
 DR PROSITE: PS50058; G\_PROTEIN\_GAMMA; 1.  
 KW Transducer; Prenylation; Lipoprotein; Multigene family; 3D-structure.  
 FT INIT MET 0 0  
 FT DISULFID 35 36 FARNESYL.  
 FT LIPID 70 70 REMOVED IN MATURE FORM.  
 FT PROPEP 71 73 B1743E36F4BD2505 CRC64;  
 SQ SEQUENCE 73 AA; 8413 MW;  
 Query Match 18.2%; Score 36; DB 1; Length 73;  
 Best Local Similarity 42.1%; Pred. No. 1.9e+02;  
 Matches 16; Conservative 2; Mismatches 10; Indels 10; Gaps 4;  
 Oy 3 QNFRVYV--RDSRPLPMKG-----PAKLWKGCAV 32  
 Db 37 EEFRDYVERSGEDPLVKIPEDKPKFKEL-KG-GCVI 72  
 RESULT 7  
 R35A\_PVRAB STANDARD; PRT; 87 AA.  
 ID R35A\_PVRAB  
 AC Q9V1P2;  
 DT 16-OCT-2001 (Rel. 40, Created)  
 DT 16-OCT-2001 (Rel. 40, Last sequence update)  
 DT 16-OCT-2001 (Rel. 40, Last annotation update)  
 DE 50S ribosomal protein L35ae.  
 GN RPL35AE OR PAB7092.  
 OS Pyrococcus abyssi.  
 OC Archaea; Euryarchaeota; Thermococci; Thermococcales; Thermococcaceae;  
 CC NCBI\_TaxID=29292;  
 RN [1]  
 RP SEQUENCE FROM N.A.  
 RC STRAIN=GES / Orsay;  
 RA Heilig R.;  
 RT "Pyrococcus abyssi genome sequence: insights into archaeal chromosome  
 RT structure and evolution.";  
 RL Submitted (JUL-1999) to the EMBL/GenBank/DBJ databases.  
 [16]  
 RP SIMILARITY: BELONGS TO THE L35AE FAMILY OF RIBOSOMAL PROTEINS.  
 [17]  
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 CC -----  
 DR EMBL: AJ248284; CAB49307.1; -  
 DR InterPro: IPR001780; Ribosomal\_L35AE.  
 DR Pfam: PF01247; Ribosomal\_L35ae; 1.  
 DR ProDom: PD012670; Ribosomal\_L35AE; 1.  
 DR PROSITE: PS01105; RIBOSOMAL\_L35AE; 1.  
 KW Ribosomal protein; Complete proteome.  
 SQ SEQUENCE 87 AA; 9736 MW; 79A625A3AF744958 CRC64;

Query Match 18.2%; Score 36; DB 1; Length 87;  
Best Local Similarity 46.2%; Pred. No. 2.3e+02;  
Matches 6; Conservative 3; Mismatches 4; Indels 0; Gaps 0;

Oy 16 LMKGPAKLKMG 28  
Db 44 LMKSPGKILKMG 56

## RESULT 8

R35A\_PYRFU STANDARD; PRT; 87 AA.  
AC Q8T2V6;  
DT 15-JUN-2002 (Rel. 41, Created)  
DT 15-JUN-2002 (Rel. 41, Last sequence update)  
DE 508 ribosomal protein L35Ae.  
GN RPL35AE OR PFI872.  
OS Pyrococcus furiosus.  
OC Archaea; Euryarchaeota; Thermococci; Thermococcales; Thermococcaceae;  
OX NCBI\_TaxID=2261;  
RN [1]  
RP SEQUENCE FROM N.A.  
RC STRAIN=Vci / DSM 3638 / ATCC 43587 / JCM 8422;  
RA Weiss R.B., Dunn D.M., Robb F.T., Brown J.R.;  
RT "The complete sequence of the Pyrococcus furiosus genome."  
RL Submitted (FEB-2002) to the EMBL/GenBank/DBJ databases.  
CC -1- SIMILARITY: BELONGS TO THE L35AE FAMILY OF RIBOSOMAL PROTEINS.  
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DR EMBL; AE010282; AAL81996.1.  
DR InterPro; IPR001780; Ribosomal\_L35AE.  
DR Pfam; PF01247; Ribosomal\_L35AE; 1.  
DR ProDom; PD012670; Ribosomal\_L35AE; 1.  
DR PROSITE; PS01105; RIBOSOMAL\_L35AE; 1.  
KW Ribosomal protein; Complete proteome.  
SQ SEQUENCE 87 AA; 9735 MW; FSC425AA3A09247 CRC64;

Query Match 18.2%; Score 36; DB 1; Length 87;  
Best Local Similarity 46.2%; Pred. No. 2.3e+02;  
Matches 6; Conservative 3; Mismatches 4; Indels 0; Gaps 0;

Oy 16 LMKGPAKLKMG 28  
Db 44 LMKSPGKILKMG 56

## RESULT 9

R35A\_PYRFU STANDARD; PRT; 87 AA.  
AC P20299;  
DT 01-FEB-1991 (Rel. 17, Created)  
DT 01-FEB-1991 (Rel. 17, Last sequence update)  
DT 30-MAY-2000 (Rel. 39, Last annotation update)  
DE 508 ribosomal protein L35Ae.  
GN RPL35AE.  
OS Pyrococcus woessli.  
OC Archaea; Euryarchaeota; Thermococci; Thermococcales; Thermococcaceae;  
OX NCBI\_TaxID=2262;  
RN [1]  
RP SEQUENCE FROM N.A.  
RC STRAIN=DSM 3773;  
RX MEDLINE=90330536; PubMed=2165475;

RA Zwickl P., Fabry S., Bogedain C., Haas A., Hensel R.;  
RT "Glyceroldehyde-3-phosphate dehydrogenase from the hyperthermophilic  
RT archaeobacterium Pyrococcus woessli: characterization of the enzyme,  
RT cloning and sequencing of the gene, and expression in Escherichia  
RT coli."  
RL J. Bacteriol. 172:4329-4338(1990).

CC -1- SIMILARITY: BELONGS TO THE L35AE FAMILY OF RIBOSOMAL PROTEINS.  
DR PIR; S10651; OOOYVW.  
DR InterPro; IPR001780; Ribosomal\_L35AE.  
DR Pfam; PF01247; Ribosomal\_L35AE; 1.  
DR ProDom; PD012670; Ribosomal\_L35AE; 1.  
DR PROSITE; PS01105; RIBOSOMAL\_L35AE; 1.  
KW Ribosomal protein.  
SQ SEQUENCE 87 AA; 9666 MW; B5D524BBA3A148EC CRC64;

Query Match 18.2%; Score 36; DB 1; Length 87;  
Best Local Similarity 46.2%; Pred. No. 2.3e+02;  
Matches 6; Conservative 3; Mismatches 4; Indels 0; Gaps 0;

Oy 16 LMKGPAKLKMG 28  
Db 44 LMKSPGKILKMG 56

## RESULT 10

YVAP\_VACCC STANDARD; PRT; 89 AA.  
AC P20525;  
DT 01-FEB-1991 (Rel. 17, Created)  
DT 01-FEB-1991 (Rel. 17, Last sequence update)  
DT 16-OCT-2001 (Rel. 40, Last annotation update)  
DE Hypothetical 9.9 kDa protein.  
GN A ORF P. protein (strain Copenhagen).

OS Vaccinia virus (strain Copenhagen).  
OC Viruses; dsDNA viruses, no RNA stage; Poxviridae; Chordopoxvirinae;  
OC Orthopoxvirus.  
OX NCBI\_TaxID=10249;  
RN [1]  
RP SEQUENCE FROM N.A.  
RX MEDLINE=91021027; PubMed=2219722;  
RA Goebel S.J., Johnson G.P., Perkins M.E., Davis S.W., Winslow J.P.,  
RA Paoletti E.;  
RT "The complete DNA sequence of vaccinia virus."  
RL Virology 179:247-266(1990).

RN [2]  
RP COMPLETE GENOME.  
RA Goebel S.J., Johnson G.P., Perkins M.E., Davis S.W., Winslow J.P.,  
RA Paoletti E.;  
RT "Appendix to 'The complete DNA sequence of vaccinia virus'";  
RL Virology 179:517-563(1990).

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DR EMBL; M35027; AAA48170.1.  
DR PIR; C42525; C42525.  
KW Hypothetical protein.  
SQ SEQUENCE 89 AA; 9909 MW; 399EA9270DF3E75A CRC64;

Query Match 18.2%; Score 36; DB 1; Length 89;  
Best Local Similarity 35.7%; Pred. No. 2.4e+02;  
Matches 5; Conservative 5; Mismatches 4; Indels 0; Gaps 0;

Oy 19 GPAKLKMGAGAV 32  
Db 24 GPSKIGWLKGFII 37

RESULT 11  
BARS\_BACAM STANDARD; PRT; 89 AA.  
ID P11540;  
AC 01-OCT-1989 (Rel. 12, Created)  
DT 01-JAN-1990 (Rel. 13, Last sequence update)  
DE 15-JUL-1999 (Rel. 38, Last annotation update)  
DR Barsear (Ribonuclease inhibitor).  
OS Bacillus amyloliquefaciens.  
OC Bacteria; Firmicutes; Bacillales; Bacillaceae; Bacillus.  
NCBI\_TaxId=1390;  
[1]  
RN SEQUENCE FROM N.A.  
RX MEDLINE=99012012; PubMed=3050134;  
RA Hartley R.W.;  
RT "Barnase and barsear. Expression of its cloned inhibitor permits  
expression of a cloned ribonuclease.";  
J. Mol. Biol. 202:913-915(1988).  
[2]  
RP REVIEW.  
RX MEDLINE=90162921; PubMed=2696173;  
RA Hartley R.W.;  
RT "Barnase and Barsear: two small proteins to fold and fit together.";  
Trends Biochem. Sci. 14:450-454(1989).  
[3]  
RP X-RAY CRYSTALLOGRAPHY (2.6 ANGSTROMS) OF COMPLEX WITH BARNASE.  
RA Guillet V., Lapthorn A., Hartley R.W., Mauguen Y.;  
RT "Recognition between a bacterial ribonuclease, barnase, and its  
natural inhibitor, Barsear.";  
Structure 1:165-177(1993).  
[4]  
RP X-RAY CRYSTALLOGRAPHY (1.7 ANGSTROMS) OF COMPLEX WITH RNASE SA.  
RX MEDLINE=96437624; PubMed=9757110;  
RA Sevcik J., Urbanikova L., Dauter Z., Wilson K.S.;  
RT "Recognition of RNase Sa by the inhibitor barsear: structure of the  
complex at 1.7 A resolution.";  
Acta Crystallogr. D 54:954-963(1998).  
[5]  
RP STRUCTURE BY NMR.  
RX MEDLINE=94009694; PubMed=8405454;  
RA Ludlenski M.J., Bycroft M., Jones D.N.M., Fersht A.R.;  
RT "Assignment of the backbone 1H and 15N NMR resonances and secondary  
structure characterization of Barsear.";  
FEBS Lett. 332:81-87(1993).  
[6]  
RP STRUCTURE BY NMR.  
RX MEDLINE=94318630; PubMed=8043574;  
RA Ludlenski M.J., Bycroft M., Freund S.M.V., Fersht A.R.;  
RT "Three-dimensional solution structure and 13C assignments of Barsear  
using nuclear magnetic resonance spectroscopy.";  
Biochemistry 33:8866-8877(1994).  
-1- FUNCTION: INHIBITOR OF THE RIBONUCLEASE BARNASE. FORMS A ONE-TO-  
ONE NON-COVALENT COMPLEX.  
-1- SUBCELLULAR LOCATION: Cytoplasmic.  
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-----  
DR EMBL; X15345; CAA33551.1; -  
DR PIR; S01373; S01373.  
DR PDB; 1BRS; 31-JUL-94.  
DR PDB; 1BTA; 31-JUL-94.  
DR PDB; 1BTB; 04-JUL-94.  
DR PDB; 1AB7; 04-SEP-97.  
DR PDB; 1A19; 08-APR-98.  
DR PDB; 1B27; 09-DEC-98.  
DR PDB; 1B25; 09-DEC-98.  
DR PDB; 1B20; 09-DEC-98.

DR PDB; 1B35; 09-DEC-98.  
DR PDB; 1AV7; 02-MAR-99.  
DR InterPro; IPR000468; Barsear.  
DR Pfam; PF01337; Barsear; 1.  
DR ProDom; PD029050; Barsear; 1.  
KW 3D-structure.  
FT INIT MET 0  
SQ SEQUENCE 89 AA; 10212 MW; 3AC7E76A9C43A505 CRC64;  
Query Match 17.9%; Score 35.5; DB 1; Length 89;  
Best Local Similarity 30.8%; Pred. No. 2.8e+02;  
Matches 8; Conservative 4; Mismatches 7; Indels 7; Gaps 1;  
OY 8 YYPDSRPDM-----KSPAKLWK 26  
DB 29 YIGENDALMDCLTGMVEPLVLEWR 54

RESULT 12  
SR19\_METH STANDARD; PRT; 91 AA.  
ID AC 026267;  
DT 16-OCT-2001 (Rel. 40, Created)  
DT 16-OCT-2001 (Rel. 40, Last sequence update)  
DE Signal recognition particle 19 kDa protein (SRP19).  
GN SRP19 OR MTH165.  
OS Mechanobacterium thermoautotrophicum.  
OC Archaea; Euryarchaeota; Methanobacteria; Methanobacteriales;  
OC Mechanobacteriaceae; Methanothermobacter.  
CX NCBI\_TaxId=187420;  
[1]  
RP SEQUENCE FROM N.A.  
RC STRAIN=Delta H;  
RX MEDLINE=98037514; PubMed=9371463;  
RA Smith D.R., Doucette-Stamm L.A., Deloughery C., Lee H.-M., Dubois J.,  
RA Aldredge T., Bashirzadeh R., Blakely D., Cook R., Gilbert K.,  
RA Harrison D., Hoang L., Keagle P., Lumm W., Pochler B., Qiu D.,  
RA Spadefora R., Vicare R., Wang Y., Wierzbowski J., Gibson R.,  
RA Jiwani N., Caruso A., Bush D., Sater H., Patwell D., Prabhakar S.,  
RA McDougall J., Shimer G., Goyal A., Pietrovski S., Church G.M.,  
RA Daniels C.J., Mao J.-I., Rice P., Noelling J., Reeve J.N.;  
RT "Complete genome sequence of Methanobacterium thermoautotrophicum  
J. Bacteriol. 179:7135-7155(1997).  
-1- FUNCTION: SIGNAL-RECOGNITION-PARTICLE ASSEMBLY, BINDS DIRECTLY TO  
7S RNA AND MEDIATES BINDING OF THE 54 KDA SUBUNIT OF THE SRP (BY  
SIMILARITY).  
CC -1- SUBUNIT: ARCHAEL. SIGNAL RECOGNITION PARTICLE CONSISTS OF A 7S RNA  
MOLECULE OF 300 NUCLEOTIDES AND TWO PROTEIN SUBUNITS: SRP54 AND  
SRP19 (BY SIMILARITY).  
CC -1- SUBCELLULAR LOCATION: Cytoplasmic (By similarity).  
CC -1- SIMILARITY: BELONGS TO THE SRP19 FAMILY.  
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-----  
DR EMBL; AE000804; AAB84671.1; -  
DR InterPro; IPR002778; SRP19.  
DR Pfam; PF01922; SRP19; 1.  
DR ProDom; PD006609; SRP19; 1.  
KW Signal recognition particle; RNA-binding; Ribonucleoprotein;  
KW Complete proteome.  
SQ SEQUENCE 91 AA; 10535 MW; BFD64D1A20B48141 CRC64;  
Query Match 17.9%; Score 35.5; DB 1; Length 91;  
Best Local Similarity 31.8%; Pred. No. 2.9e+02;  
Matches 7; Conservative 3; Mismatches 5; Indels 7; Gaps 1;

Qy 15 PLMKGPATLWKSGAVVIOIN 36  
 Db 57 PSW-----WESSGRVVEYN 71

RESULT 13  
 A4\_MACMU STANDARD; PRT; 76 AA.

AC P28216;  
 DT 01-DEC-1992 (Rel. 24, Created)  
 DT 01-DEC-1992 (Rel. 24, Last sequence update)  
 DT 16-OCT-2001 (Rel. 40, Last annotation update)  
 DE Alzheimer's disease amyloid A4 protein (Fragment).

GN APP.  
 OS Macaca mulatta (Rhesus macaque).  
 OC Eukaryota; Metazoa; Chordata; Craniata; Vertebrata; Euteleostomi;  
 OC Mammalia; Eutheria; Primates; Catarrhini; Cercopithecoidea;  
 OC Cercopithecinae; Macaca.  
 NC NCB1\_TaxID=9544;  
 RN [1]

SEQUENCE FROM N.A.

RC TISSUE=Brain;

RA Koo E.H., Sisodia S.S., Price D.L.;

RL Submitted (JUL-1989) to the EMBL/Genbank/DBJ databases.

CC -1- FUNCTION: FUNCTIONAL NEURONAL RECEPTOR WHICH COUPLES TO

CC INTRACELLULAR SIGNALING PATHWAY THROUGH THE GTP-BINDING PROTEIN

CC G(O) (BY SIMILARITY).

CC -1- SUBCELLULAR LOCATION: Type I membrane protein.

CC -1- ALTERNATIVE PRODUCTS: 5 ISOFORMS; APP(395), APP(563), APP(695),

CC APP(751) AND APP(770) (SHOWN HERE); ARE PRODUCED BY ALTERNATIVE

CC SPLICING (BY SIMILARITY).

CC -1- SIMILARITY: BELONGS TO THE APP FAMILY.

CC -1- SIMILARITY: CONTAINS 1 BPTI/KUNITZ INHIBITOR DOMAIN.

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DR EMBL; X15985; CAA34116.1; -

DR PIR; S06678; S06678.

DR HSP; P05067; 1AAP.

DR InterPro; IPR001868; A4\_APP.

DR InterPro; IPR002223; Kunitz\_BPTI.

DR Pfam; PF00014; Kunitz\_BPTI; 1.

DR ProDom; PD000222; Kunitz\_BPTI; 1.

DR SMART; SM00131; KU; 1.

DR PROSITE; PS00319; A4\_EXTRA; PARTIAL.

DR PROSITE; PS00320; A4\_INTRA; PARTIAL.

DR PROSITE; PS00280; BPTI\_KUNITZ\_1; 1.

DR PROSITE; PS50279; BPTI\_KUNITZ\_2; 1.

KW Glycoprotein; Amyloid; Neurone; Alternative splicing;

KW Serine protease inhibitor.

FT DOMAIN 1 1

FT DONTER 1 1

FT ACT\_SITE 13 14 BPTI/KUNITZ INHIBITOR.

FT DISULFID 3 53 REACTIVE BOND.

FT DISULFID 12 36 BY SIMILARITY.

FT DISULFID 28 49 BY SIMILARITY.

FT NON\_TER 76 76

SEQUENCE 76 AA; 8527 MW; 492BPF3069AB082A1 CRC64;

Query Match 17.7%; Score 35; DB 1; Length 76;

Best Local Similarity 42.9%; Pred. No. 2.8e+02;

Matches 6; Conservative 4; Mismatches 4; Indels 0; Gaps 0;

Qy 10 RDSRDPMLWGPATL 23

Db 63 KTRRPLTRDRPVKL 76

RESULT 14  
 VG7\_SpVIR STANDARD; PRT; 83 AA.

AC P15898;  
 DT 01-APR-1990 (Rel. 14, Created)  
 DT 01-FEB-1996 (Rel. 33, Last sequence update)  
 DT 01-FEB-1996 (Rel. 33, Last annotation update)  
 DE Gene 7 protein.

GN 7.

OS Spiroplasma virus SpV1-R8A2 B.

OC Viruses; ssDNA viruses; Inoviridae; Plectrovirus.

NC NCB1\_TaxID=10854;

RN [1]

SEQUENCE FROM N.A.

RA MEDLINE=90206799; PubMed=2320423;

RA Renaudin J., Aulio P., Vignault J.C., Bove J.M.;

RT "Complete nucleotide sequence of the genome of Spiroplasma citri

virus SpV1-R8A2 B.";

RL Nucleic Acids Res. 18:1293-1293(1990).

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CC -----

DR EMBL; X51344; CAA35731.1; -

DR PIR; S31018; S31018.

DR HSP; P05067; 1AAP.

DR InterPro; IPR001868; A4\_APP.

DR InterPro; IPR002223; Kunitz\_BPTI.

DR Pfam; PF00014; Kunitz\_BPTI; 1.

DR ProDom; PD000222; Kunitz\_BPTI; 1.

DR SMART; SM00131; KU; 1.

DR PROSITE; PS00319; A4\_EXTRA; PARTIAL.

DR PROSITE; PS00320; A4\_INTRA; PARTIAL.

DR PROSITE; PS00280; BPTI\_KUNITZ\_1; 1.

DR PROSITE; PS50279; BPTI\_KUNITZ\_2; 1.

KW Glycoprotein; Amyloid; Neurone; Alternative splicing;

KW Serine protease inhibitor.

FT DOMAIN 1 1

FT DONTER 1 1

FT ACT\_SITE 13 14 BPTI/KUNITZ INHIBITOR.

FT DISULFID 3 53 REACTIVE BOND.

FT DISULFID 12 36 BY SIMILARITY.

FT DISULFID 28 49 BY SIMILARITY.

FT NON\_TER 76 76

SEQUENCE 76 AA; 8527 MW; 492BPF3069AB082A1 CRC64;

Query Match 17.7%; Score 35; DB 1; Length 83;

Best Local Similarity 45.5%; Pred. No. 3.1e+02;

Matches 5; Conservative 2; Mismatches 4; Indels 0; Gaps 0;

Qy 16 LMKGPATLWK 26

Db 28 IWTGUSALWK 38

RESULT 15  
 VG73\_BPTL5 STANDARD; PRT; 85 AA.

AC Q05288;  
 DT 01-FEB-1994 (Rel. 28, Created)  
 DT 01-FEB-1994 (Rel. 28, Last sequence update)  
 DT 01-FEB-1994 (Rel. 28, Last annotation update)  
 DE Gene 73 protein (GP73).

GN 73.

OS Mycobacteriophage L5.

OC Viruses; dsDNA viruses, no RNA stage; Caudovirales; Siphoviridae;

OC L5-like viruses.

NC NCB1\_TaxID=31757;

RN [1]

SEQUENCE FROM N.A.

RA MEDLINE=93211282; PubMed=8459766;

RA Hatfull G.F., Sarkis G.J.;

RT "DNA sequence, structure and gene expression of mycobacteriophage L5:

Mo1. Microbiol. 7:395-405(1993).

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CC -----

DR EMBL; Z18946; CAA79449.1; -

DR PIR; S31018; S31018.

SQ SEQUENCE 85 AA; 10250 MM; 5032B5A1400FF4A7 CRC64;  
 Query Match 17.7%; Score 35; DB 1; Length 85;  
 Best Local Similarity 32.4%; Pred. No. 3.2e+02;  
 Matches 11; Conservative 4; Mismatches 13; Indels 6; Gaps 2;  
 QY 2 IQFRVYRDSRDEPLMKGPAPAKLWKGEGAVVIO 35  
 Db 28 IKDIERKWFADHDQ-WK-----TWHDPEAPFEQD 55

Search completed: May 6, 2003, 14:57:41  
 Job time : 12 secs



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OM protein - protein search, using sw model

Run on: May 6, 2003, 14:56:24 ; Search time 16 seconds  
(without alignments)  
216.302 Million cell updates/sec

Title: US-09-868-399-1

Perfect score: 198  
Sequence: 1 KIQNFRVYRDSRDP LMKGPALKLMKGGA VVIQDN 36

Scoring table: BLOSUM62  
Gapop 10.0, Gapext 0.5

Searched: 283224 seqs, 96134422 residues

Total number of hits satisfying chosen parameters: 37666

Minimum DB seq length: 0  
Maximum DB seq length: 100

Post-processing: Minimum Match 0%  
Maximum Match 100%

Listing first 45 summaries

Database :  
1: PIR.73:\*  
2: PIR1:\*  
3: PIR3:\*  
4: PIR4:\*

Pred. No. is the number of results predicted by chance to have a score greater than or equal to the score of the result being printed, and is derived by analysis of the total score distribution.

#### SUMMARIES

Result No.	Score	Query Match	Length	ID	Description
1	53.5	27.0	70	2 A69842	hypothetical prote
2	50.5	25.5	91	2 A97004	barstar-like prote
3	50	25.3	44	2 T09381	pol polypotein -
4	48.5	24.5	95	2 A81176	ribonuclease inhib
5	45	22.7	82	2 B95846	hypothetical prote
6	42	21.2	76	2 AD2538	hypothetical prote
7	41.5	21.0	96	2 G83600	hypothetical prote
8	41	20.7	94	2 A69848	hypothetical prote
9	40.5	20.5	68	2 AG3217	hypothetical prote
10	39.5	19.9	63	2 S26796	ig heavy chain V r
11	39.5	19.9	87	2 T03193	hypothetical prote
12	39.5	19.9	88	2 T36458	hypothetical prote
13	39.5	19.9	96	2 T36605	hypothetical prote
14	39	19.7	62	2 A12304	hypothetical prote
15	39	19.7	64	2 T05933	probable 3-methyl-
16	39	19.7	79	2 T37100	hypothetical prote
17	39	19.7	96	2 C64900	outer membrane por
18	38.5	19.4	61	2 C75321	preprotein translo
19	38.5	19.4	62	2 T26847	hypothetical prote
20	38.5	19.4	66	2 S32027	Sp12 protein homol
21	38.5	19.4	96	2 A71151	hypothetical prote
22	38.5	19.4	96	2 S75031	hypothetical prote
23	38	19.2	60	2 H81170	conserved hypotet
24	38	19.2	81	2 B95286	hypothetical prote
25	38	19.2	83	2 C72800	gpi protein - Myco
26	38	19.2	96	2 H81147	hypothetical prote
27	38	19.2	100	2 F95369	hypothetical prote
28	37.5	18.9	60	2 H97179	hypothetical prote
29	37.5	18.9	67	2 A71915	CB/ELIP/HLIP supe

30	37.5	18.9	73	2 I39159	GTP-binding regula
31	37.5	18.9	88	2 T17532	hypothetical prote
32	37	18.7	50	2 S68843	sucrose 1P-fructos
33	37	18.7	56	2 B69185	hypothetical prote
34	37	18.7	76	2 B97816	hypothetical prote
35	37	18.7	77	2 A83468	hypothetical prote
36	37	18.7	82	2 H84046	hypothetical prote
37	37	18.7	94	2 H84041	hypothetical prote
38	36.5	18.4	62	2 D69045	hypothetical prote
39	36.5	18.4	88	2 A69347	translational initia
40	36.5	18.4	97	2 E90278	hypothetical prote
41	36.5	18.4	98	2 F90314	hypothetical prote
42	36.5	18.4	99	2 E90338	hypothetical prote
43	36	18.2	36	2 C64039	hypothetical prote
44	36	18.2	46	2 F87527	hypothetical prote
45	36	18.2	52	2 S21185	beta-fructofuranos

#### ALIGNMENTS

RESULT 1  
A69842  
hypothetical protein ylx - Bacillus subtilis  
C:Species: Bacillus subtilis  
C:Date: 05-Dec-1997 #sequence\_revision 05-Dec-1997 #text\_change 15-Oct-1999  
C:Accession: A69842  
R:Kunze, F.; Ogasawara, N.; Moszer, I.; Albertini, A.M.; Alloni, G.; Aevedo, V.; Berte  
C.; Bron, S.; Brouillet, S.; Bruschi, C.V.; Caldwell, B.; Capuano, V.; Carter, N.M.; Ch  
A.; Ehrlich, S.D.; Emerson, P.T.; Entian, K.D.; Errington, J.; Fabret, C.; Ferrari, E.  
Nature 390, 249-256, 1997  
A:Authors: Foulger, D.; Fritz, C.; Fujita, M.; Fujita, Y.; Fuma, S.; Galizzi, A.; Gall  
iech, J.; Harwood, C.R.; Henaut, A.; Hilbert, H.; Holsappel, S.; Hosono, S.; Hullo, M.F  
Koetter, P.; Koningsstein, G.; Krogh, S.; Kumano, M.; Kurita, K.; Lapidus, A.; Lardinois  
A:Authors: Lauber, J.; Lazarevic, V.; Lee, S.M.; Levine, A.; Liu, H.; Masuda, S.; Maue  
Y, M.; Ogawa, K.; Ogiwara, A.; Oudega, B.; Park, S.H.; Parro, V.; Pohl, T.M.; Portetelli  
Rieger, M.; Rivolta, C.; Roche, E.; Roche, B.; Rose, M.; Sadale, Y.; Sato, T.; Scanlon  
A:Authors: Schleicher, S.; Schroeter, R.; Scofield, F.; Sekiguchi, J.; Sekowska, A.; Sero  
akeuchi, M.; Tamakoshi, A.; Tanaka, T.; Terresta, P.; Tognoni, A.; Tosato, V.; Uchiyama  
T.; Winters, P.; Wipat, A.; Yamamoto, H.; Yamane, K.; Yasumoto, K.; Yata, K.; Yoshida  
A:Authors: Yoshikawa, H.F.; Zumbstein, E.; Yoshikawa, H.; Danchin, A.  
A:Title: The complete genome sequence of the Gram-positive bacterium Bacillus subtilis.  
A:Reference number: A69580; MUID:98044033; PMID:9384377  
A:Accession: A69842  
A:Status: preliminary; nucleic acid sequence not shown; translation not shown  
A:Residues: 1-70 <KUN>  
A:Cross-references: GB:299109; GB:AL009126; NID:g2633260; PIDN:CA812957.1; PID:e1183119  
A:Experimental source: strain 168  
C:Genetics:  
A:Gene: ylx  
Query Match 27.0%; Score 53.5; DB 2; Length 70;  
Best Local Similarity 42.9%; Pred. No. 1.8;  
Matches 12; Conservative 5; Mismatches 10; Indels 1; Gaps 1;  
Cy 6 RYVYRDSRDP LMKGPALKLMKGGA VVI 33  
Db 44 RLVYR-RPPCKRTGKXIMWEDAVV 70  
RESULT 2  
A97004  
barstar-like protein ribonuclease (barnase) inhibitor [imported] - Clostridium acetobut  
C:Species: Clostridium acetobutylicum  
C:Date: 14-Sep-2001 #sequence\_revision 14-Sep-2001 #text\_change 14-Sep-2001  
C:Accession: A97004  
R:Neilling, J.; Bretton, G.; Omelchenko, M.V.; Markarova, K.S.; Zeng, Q.; Gibson, R.; Lee  
J.; Daly, M.J.; Bennett, G.N.; Koonin, E.V.; Smith, D.R.  
J. Bacteriol. 183, 4823-4838, 2001  
A:Title: Genome Sequence and Comparative Analysis of the Solvent-Producing Bacterium C1  
A:Reference number: A96900; MUID:21359325; PMID:21359325  
A:Accession: A97004

A/Status: preliminary  
 A/Molecule type: DNA  
 A/Residues: 1-91 <KUR>  
 A/Cross-references: GB:AE001437; PIDN:AAK78820.1; PID:915023737; GSPDB:GN00168  
 A/Experimental source: Clostridium acetobutylicum ATCC824  
 C/Genetics:  
 A/Gene: CAC0844

Query Match 25.5%; Score 50.5; DB 2; Length 91;  
 Best Local Similarity 37.9%; Pred. No. 6.2;  
 Matches 11; Conservative 3; Mismatches 8; Indels 7; Gaps 1;

QY 5 FRYVYDSDPDM-----KCAPLTK 26  
 DB 27 FRYVYDSDPDM-----KCAPLTK 55

## RESULT 3

109381  
 pol polyprotein - human immunodeficiency virus type 1 (isolate cntrl 1) (fragment)  
 C/Species: human immunodeficiency virus type 1, HIV-1  
 A/Variety: isolate cntrl 1  
 C/Date: 11-Jun-1999 #sequence\_revision 11-Jun-1999 #text\_change 23-Jul-1999  
 C/Accession: T09381

R/McCosh, N.L.; Chang, G.; d'Arcy, L.A.; Ehrenberg, P.K.; Mariani, R.; Busch, M.P.; Bir  
 J. Virol. 69, 4228-4236, 1995  
 A/Title: Defective accessory genes in a human immunodeficiency virus type 1-infected lo  
 A/Reference number: Z16554; MUID:95287475; PMID:7769682  
 A/Accession: T09381  
 A/Status: preliminary; translated from GB/EMBL/DBJ  
 A/Molecule type: DNA  
 A/Residues: 1-44 <MTC>  
 A/Cross-references: EMBL:U24451; NID:9829440; PID:9829441  
 C/Genetics:  
 A/Gene: pol  
 C/Superfamily: pol polyprotein  
 C/Keywords: AIDS; immunodeficiency; polyprotein

Query Match 25.3%; Score 50; DB 2; Length 44;  
 Best Local Similarity 100.0%; Pred. No. 3.2;  
 Matches 10; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

QY 27 GEGAVIODN 36  
 DB 1 GEGAVIODN 10

## RESULT 4

A81176  
 ribonuclease inhibitor barstar NMB0646 [imported] - Neisseria meningitidis (strain MCS8  
 C/Species: Neisseria meningitidis  
 C/Date: 31-Mar-2000 #sequence\_revision 31-Mar-2000 #text\_change 19-Jan-2001  
 C/Accession: A81176

R/Tettelin, H.; Saunders, N.J.; Heidelberg, J.; Jeffries, A.C.; Nelson, K.E.; Eisen, J.A  
 Hickey, E.K.; Haft, D.H.; Salzberg, S.L.; White, O.; Fleischmann, R.D.; Dougherty, B.A.;  
 ri, H.; Qin, H.; Vamathevan, J.; Gill, J.; Scarlato, V.; Masiiani, V.; Piza, M.  
 Science 287, 1809-1815, 2000  
 A/Authors: Grandi, G.; Sun, L.; Smith, H.O.; Fraser, C.M.; Moxon, E.R.; Rappuoli, R.; Ve  
 A/Title: Complete genome sequence of Neisseria meningitidis serogroup B strain MCS8..  
 A/Reference number: A81000; MUID:2017575; PMID:10710307  
 A/Accession: A81176  
 A/Status: preliminary  
 A/Molecule type: DNA  
 A/Residues: 1-95 <TET>  
 A/Cross-references: GB:AE002419; GB:AE002098; NID:97225863; PIDN:AAF1067.1; PID:9722587  
 A/Experimental source: serogroup B, strain MCS8  
 C/Genetics:  
 A/Gene: NMB0646

Query Match 24.5%; Score 48.5; DB 2; Length 95;  
 Best Local Similarity 33.3%; Pred. No. 12;  
 Matches 11; Conservative 6; Mismatches 9; Indels 7; Gaps 1;

QY 1 KIONFRVYDSDPDM-----KCAPLTK 26  
 DB 22 KIFSTIDYGNMIDALMDLSTVVERPITLVK 54

## RESULT 5

B95846  
 hypothetical protein [imported] - Sinorhizobium meliloti (strain 1021) megaplasmid pSym  
 C/Species: Sinorhizobium meliloti  
 C/Date: 24-Aug-2001 #sequence\_revision 24-Aug-2001 #text\_change 30-Sep-2001  
 C/Accession: B95846

R/Finan, T.M.; Weidner, S.; Wong, K.; Bunhmaster, J.; Chain, P.; Vorholter, F.J.; Herra  
 Proc. Natl. Acad. Sci. U.S.A. 98, 9889-9894, 2001  
 A/Title: The complete sequence of the 1,683-kb pSymB megaplasmid from the N2-fixing en  
 A/Reference number: A95842; MUID:21396508; PMID:11481431  
 A/Accession: B95846  
 A/Status: preliminary  
 A/Molecule type: DNA  
 A/Residues: 1-82 <KUR>  
 A/Cross-references: GB:AL591985; PIDN:CAC48434.1; PID:915139906; GSPDB:GN00167  
 A/Experimental source: strain 1021, megaplasmid pSymB  
 R/Galibert, F.; Finan, T.M.; Long, S.R.; Puhler, A.; Abola, P.; Ampe, F.; Barloy-Hubler  
 eta, D.; Chain, P.; Cowie, A.; Davis, R.W.; Dreano, S.; Federspiel, N.A.; Fisher, R.F  
 Science 293, 668-672, 2001  
 A/Authors: Kahn, D.; Kahn, M.L.; Keating, D.H.; Kisse, E.; Komp, C.; Lelaure  
 hebaull, P.; Vandenbol, M.; Vorholter, F.J.; Weidner, S.; Wells, D.H.; Wong, K.; Yeh, K  
 A/Title: The composite genome of the legume symbiont Sinorhizobium meliloti.  
 A/Reference number: A96039; MUID:21368234; PMID:11474104  
 C/Genetics:  
 A/Gene: SMD20031  
 A/Genome: plasmid

Query Match 22.7%; Score 45; DB 2; Length 82;  
 Best Local Similarity 37.5%; Pred. No. 32;  
 Matches 9; Conservative 4; Mismatches 11; Indels 0; Gaps 0;

QY 12 SRDPLMKGPAKLIMGEGAVIYD 35  
 DB 20 SRDGHWRDPRPLAADQIVITD 43

## RESULT 6

AD2538  
 hypothetical protein asl7591 [imported] - Nostoc sp. (strain PCC 7120) Plasmid pCC7120b.  
 C/Species: Nostoc sp.  
 A/Note: Nostoc sp. strain PCC 7120 is a synonym of Anabaena sp. strain PCC 7120  
 C/Date: 14-Dec-2001 #sequence\_revision 14-Dec-2001 #text\_change 30-Jun-2002  
 C/Accession: AD2538

R/Kaneke, T.; Nakamura, Y.; Wolk, C.P.; Kuritz, T.; Sasaoto, S.; Watanabe, A.; Iriyuchi  
 Nakazaki, N.; Shimpo, S.; Sugimoto, M.; Takazawa, M.; Yamada, M.; Yasuda, M.; Tabata, S  
 DNA Res. 8, 205-213, 2001  
 A/Title: Complete genomic sequence of the filamentous Nitrogen-fixing Cyanobacterium An  
 A/Reference number: AB1807; MUID:21595285; PMID:11759640  
 A/Accession: AD2538  
 A/Status: preliminary  
 A/Molecule type: DNA  
 A/Residues: 1-76 <KUR>  
 A/Cross-references: GB:AP003602; PIDN:BAF7234.1; PID:917134676; GSPDB:GN00181  
 A/Experimental source: strain PCC 7120  
 C/Genetics:  
 A/Gene: asl7591  
 A/Genome: plasmid

Query Match 21.2%; Score 42; DB 2; Length 76;  
 Best Local Similarity 40.0%; Pred. No. 78;  
 Matches 6; Conservative 5; Mismatches 4; Indels 0; Gaps 0;

QY 9 YDSDPDMKGPATL 23  
 DB 3 HOSDSDSWWRSPAKI 17

RESULT 7  
G83600  
hypothetical protein PA0369 [imported] - *Pseudomonas aeruginosa* (strain PA01)  
C/Species: *Pseudomonas aeruginosa*  
C/Date: 15-Sep-2000 #sequence\_revision 15-Sep-2000 #text\_change 31-Dec-2000  
C/Accession: G83600  
R/Owner: C.K.; Pham, X.Q.; Eryin, A.L.; Mizoguchi, S.D.; Warener, P.; Hickey, M.J.; Br  
adman, S.; Yuan, Y.; Brody, L.L.; Coulter, S.N.; Folger, K.R.; Kas, A.; Lardis, K.; Lim,  
L.; Lo, S.; Olson, M.V.  
Nature 406, 959-964, 2000  
A/Title: Complete genome sequence of *Pseudomonas aeruginosa* PA01, an opportunistic patho  
gen  
A/Reference number: A82950; MUID:20437337; PMID:10984043  
A/Accession: G83600  
A/Status: preliminary  
A/Molecule type: DNA  
A/Residues: 1-96 <STO>  
A/Cross-references: GB:AE004474; GB:AE004091; NID:G9946210; PIDN:AG03758.1; GSPDB:GN001  
A/Experimental source: strain PA01  
C/Genetics:  
A/Gene: PA0369

Query Match 21.0%; Score 41.5; DB 2; Length 96;  
Best Local Similarity 32.1%; Pred. No. 1.2e+02;  
Matches 9; Conservative 5; Mismatches 9; Indels 5; Gaps 1;

OY 14 DRLMKPAKL-----LWKEGAVVQDN 36  
DB 35 DDLWQMAASLACFGWWSGAGEDILDD 62

RESULT 8  
A69848  
hypothetical protein yjC0 - *Bacillus subtilis*  
C/Species: *Bacillus subtilis*  
C/Date: 05-Dec-1997 #sequence\_revision 05-Dec-1997 #text\_change 15-Oct-1999  
C/Accession: A69848  
R/Kunst, F.; Ogasawara, N.; Moszer, I.; Albertini, A.M.; Alloni, G.; Azevedo, V.; Berter  
C.; Bron, S.; Brouillet, S.; Brusch, C.V.; Caldwell, B.; Capiano, V.; Carter, N.M.; Ch  
A.; Ehrlich, S.D.; Emmerson, P.T.; Entian, K.D.; Errington, J.; Fabret, C.; Ferrari, E.  
Nature 390, 249-256, 1997  
A/Authors: Foulger, D.; Fritz, C.; Fujita, M.; Fujita, Y.; Fuma, S.; Gallazzi, A.; Gall  
Aech, J.; Harwood, C.R.; Henaut, A.; Hilbert, H.; Holstappel, S.; Hosono, S.; Hullo, M.F.  
Koeter, P.; Koningsreim, G.; Kropp, S.; Kumano, M.; Kurita, K.; Lapidus, A.; Lardinois  
A.; Authors: Lamber, J.; Lazarevic, V.; Lee, S.M.; Levine, A.; Liu, H.; Masuda, S.; Maue  
Y., M.; Ogawa, K.; Ogata, A.; Oudega, B.; Park, S.H.; Patro, V.; Sadale, Y.; Sato, T.; Scallan  
Rieger, M.; Rivolta, C.; Roche, E.; Roche, B.; Rose, M.; Sadale, Y.; Sato, T.; Scallan  
A.; Authors: Schleich, S.; Schroeter, R.; Scoffone, F.; Sekiguchi, J.; Sekowska, A.; Ser  
akeuchi, M.; Tamakoshi, A.; Tanaka, T.; Terpetra, P.; Togomi, A.; Tosato, V.; Uchiyama,  
T.; Winters, P.; Wipat, A.; Yamamoto, H.; Yamane, K.; Yasumoto, K.; Yata, K.; Yoshida, K  
A.; Authors: Yoshikawa, H.F.; Zumbstein, E.; Yoshikawa, H.; Danchin, A.  
A/Title: The complete genome sequence of the Gram-positive bacterium *Bacillus subtilis*.  
A/Reference number: A69580; MUID:98044033; PMID:9384377  
A/Accession: A69848  
A/Status: preliminary; nucleic acid sequence not shown; translation not shown  
A/Molecule type: DNA  
A/Residues: 1-94 <KIN>  
A/Cross-references: GB:Z99110; GB:AL009126; NID:G2633472; PIDN:CAB13052.1; PID:el183215;  
A/Experimental source: strain 168  
C/Genetics:  
A/Gene: yjC0

Query Match 20.7%; Score 41; DB 2; Length 94;  
Best Local Similarity 45.5%; Pred. No. 1.4e+02;  
Matches 10; Conservative 3; Mismatches 9; Indels 0; Gaps 0;

OY 7 VYRDSRDLWKGPALKMKGE 28  
DB 49 VHYSDRPHLYKGLPELTKGE 70

RESULT 9  
AG3217

hypothetical protein Atu5470 [imported] - *Agrobacterium tumefaciens* (strain C58, Dupont  
C/Species: *Agrobacterium tumefaciens*  
C/Date: 11-Jan-2002 #sequence\_revision 11-Jan-2002 #text\_change 11-Jan-2002  
C/Accession: AG3217  
R/Owner: G.D.; Setubal, J.C.; Kaul, R.; Monks, D.; Chen, L.; Wood, G.E.; Chen, Y.; Woo,  
erage, W.; Gillet, W.; Grant, C.; Guenther, D.; Kutyavina, T.; Levy, R.; Li, M.; McClel  
Karp, P.; Romero, P.; Zhang, S.  
Science 294, 2317-2323, 2001  
A/Authors: Yoo, H.; Tao, Y.; Biddle, P.; Jung, M.; Krespan, W.; Perry, M.; Gordon-Kamm,  
eter, E.W.  
A/Title: The Genome of the Natural Genetic Engineer *Agrobacterium tumefaciens* C58.  
A/Reference number: AB2577; PMID:11743193  
A/Accession: AG3217  
A/Status: preliminary  
A/Molecule type: DNA  
A/Residues: 1-68 <KUR>  
A/Cross-references: GB:AE008687; PIDN:AA146157.1; PID:G17743927; GSPDB:GN00188  
A/Experimental source: strain C58 (Dupont)  
C/Genetics:  
A/Gene: Atu5470  
A/Gene: plasmid

Query Match 20.5%; Score 40.5; DB 2; Length 68;  
Best Local Similarity 37.0%; Pred. No. 1.1e+02;  
Matches 10; Conservative 4; Mismatches 6; Indels 7; Gaps 2;

OY 9 VYRDSRDLWKGPALKMKGE 28  
DB 20 WKDIRAPLWRINGRGDQASAWAGE 46

RESULT 10  
S26796  
Ig heavy chain V region - human (fragment)  
C/Species: *Homo sapiens* (man)  
C/Date: 13-Jan-1995 #sequence\_revision 30-Apr-1999 #text\_change 20-Jun-2000  
C/Accession: S26796  
R/Mortari, F.; Newton, J.A.; Wang, J.Y.; Schroeder Jr., H.W.  
Eur. J. Immunol. 22, 241-245, 1992  
A/Title: The human cord blood antibody repertoire. Frequent usage of the V(H)7 gene fam  
A/Reference number: S26786; MUID:92111632; PMID:1730251  
A/Accession: S26796  
A/Molecule type: mRNA  
A/Residues: 1-63 <MOR>  
A/Cross-references: EMBL:X61018; NID:G32785; PIDN:CAA43352.1; PID:G1335121  
C/Superfamily: Immunoglobulin V region; Immunoglobulin homology  
C/Keywords: heterotrimer; immunoglobulin

Query Match 19.9%; Score 39.5; DB 2; Length 63;  
Best Local Similarity 31.2%; Pred. No. 1.4e+02;  
Matches 10; Conservative 9; Mismatches 8; Indels 5; Gaps 3;

OY 1 KIONFRYVYRDSRDLWKGPALKMKGEAVV 32  
DB 33 RVEDTAVVY-CARQPNWEDAP---W-GQGTLV 59

RESULT 11  
T03193  
hypothetical protein 87 - rice mitochondrion  
C/Species: mitochondrion *Oryza sativa* (rice)  
C/Date: 24-Mar-1999 #sequence\_revision 24-Mar-1999 #text\_change 21-Jul-2000  
C/Accession: T03193  
R/Adami, H.; Wakaugli, T.; Sugita, M.; Sugitara, M.; Nakazono, M.; Hirai, A.  
Plant Cell Physiol. 35, 1239-1244, 1994  
A/Title: Nucleotide sequence of a 28-kbp portion of rice mitochondrial DNA: the existen  
A/Reference number: Z14841; MUID:95211382; PMID:7545979  
A/Accession: T03193  
A/Status: preliminary; translated from GB/EMBL/DBD  
A/Molecule type: DNA  
A/Residues: 1-87 <ITA>  
A/Cross-references: EMBL:D32052; NID:G769704; PIDN:BA06814.1; PID:G769710  
A/Experimental source: cultivar Nipponbare

C/Genetics:  
A/Genome: mitochondrion  
C/Keywords: mitochondrion

Query Match 19.9%; Score 39.5; DB 2; Length 87;  
Best Local Similarity 31.0%; Pred. No. 2e+02;  
Matches 9; Conservative 7; Mismatches 12; Indels 1; Gaps 1;

QY 3 QNFRVYRDSRDLPMKGPALKLMKGEGAV 31  
DB 31 QNLI-YNORPRHCKGSKMIMIRICL 58

## RESULT 12

T36458

hypothetical protein SCP43A.35c - Streptomyces coelicolor

C/Species: Streptomyces coelicolor  
C/Date: 03-Dec-1999 #sequence\_revision 03-Dec-1999 #text\_change 03-Dec-1999

C/Accession: T36458

R/Seeger, K.; Harris, D.; James, K.D.; Parkhill, J.; Barrrell, B.G.; Rajandream, M.A.  
submitted to the EMBL Data Library, July 1999

A/Reference number: 221598

A/Accession: T36458

A/Status: preliminary; translated from GB/EMBL/DBJ

A/Molecule type: DNA

A/Residues: 1-88 &lt;SEE&gt;

A/Cross-references: EMBL:AL096837; PIDN:CAB48922.1; GSPDB:GN00070; SCOEDB:SCF43A.35c

A/Experimental source: strain A3(2)

C/Genetics:

A/Gene: SCOEDB:SCF43A.35c

## Query Match

Best Local Similarity 19.9%; Score 39.5; DB 2; Length 88;  
Matches 9; Conservative 3; Mismatches 8; Indels 1; Gaps 1;

QY 10 RDSRDLPMKGPALKLMKGGA 30  
DB 18 RDDRERLKKGD-KVTWSSHS 37

## RESULT 13

T36605

hypothetical protein SCH24.40c - Streptomyces coelicolor (fragment)

C/Species: Streptomyces coelicolor  
C/Date: 03-Dec-1999 #sequence\_revision 03-Dec-1999 #text\_change 03-Dec-1999

C/Accession: T36605

R/Oliver, K.; Harris, D.; James, K.D.; Parkhill, J.; Barrrell, B.G.; Rajandream, M.A.  
submitted to the EMBL Data Library, May 1999

A/Reference number: 221575

A/Accession: T36605

A/Status: preliminary; translated from GB/EMBL/DBJ

A/Molecule type: DNA

A/Residues: 1-96 &lt;OLI&gt;

A/Cross-references: EMBL:AL049826; PIDN:CAB42746.1; GSPDB:GN00070; SCOEDB:SCH24.40c

A/Experimental source: strain A3(2)

C/Genetics:

A/Gene: SCOEDB:SCH24.40c

## Query Match

Best Local Similarity 19.9%; Score 39.5; DB 2; Length 96;  
Matches 10; Conservative 4; Mismatches 6; Indels 1; Gaps 1;

QY 1 KIONFRVYRDSRDLPMKGPALKLMKG 20  
DB 25 KVOHFVCHLESMDPSLRHGP 45

## RESULT 14

AI2304

hypothetical protein asr3992 [imported] - Nostoc sp. (strain PCC 7120)

C/Species: Nostoc sp.

A/Note: Nostoc sp. strain PCC 7120 is a synonym of Anabaena sp. strain PCC 7120  
C/Date: 14-Dec-2001 #sequence\_revision 14-Dec-2001 #text\_change 30-Jun-2002

C/Accession: AI2304

R/Kaneko, T.; Nakamura, Y.; Wolk, C.P.; Kuritz, T.; Sasamoto, S.; Watanabe, A.; Iritugu  
Nakazaki, N.; Shimo, S.; Sugimoto, M.; Takazawa, M.; Yamada, M.; Yasuda, M.; Tabata,  
DNA Res. 8, 205-213, 2001

A/Title: Complete Genomic Sequence of the Filamentous Nitrogen-fixing Cyanobacterium An

A/Reference number: AB1807; MUID:21595285; PMID:11759840  
A/Accession: AI2304

A/Status: preliminary

A/Molecule type: DNA

A/Residues: 1-62 &lt;KUR&gt;

A/Cross-references: GB:BA000019; PIDN:BAH75691.1; PID:917133127; GSPDB:GN00179

A/Experimental source: strain PCC 7120

C/Genetics:

A/Gene: asr3992

C/Superfamily: conserved hypothetical protein ycf9

## Query Match

Best Local Similarity 19.7%; Score 39; DB 2; Length 62;  
Matches 7; Conservative 0; Mismatches 6; Indels 0; Gaps 0;

QY 17 WKGPALKLMKGE 29  
DB 33 WVESKLLMLGSG 45

## RESULT 15

T05933

probable 3-methyl-2-oxobutanoate dehydrogenase (lipoamide) (EC 1.2.4.4) alpha chain - b

N/Alternate names: branched-chain alpha-keto acid decarboxylase complex E1-alpha chain

C/Species: Hordeum vulgare (barley)

C/Date: 30-Apr-1999 #sequence\_revision 30-Apr-1999 #text\_change 20-Jun-2000

C/Accession: T05933

R/Hess, W.R.; Goiz, R.R.; Boerner, T.

Plant Sci. 133, 191-201, 1998

A/Title: Analysis of randomly selected cDNAs reveals the expression of stress- and def:

A/Reference number: Z15411

A/Accession: T05933

A/Status: preliminary

A/Molecule type: mRNA

A/Residues: 1-64 &lt;HES&gt;

A/Cross-references: EMBL:AJ222787; PIDN:CAI10992.1

A/Experimental source: cv. Haisa, leaf

C/Superfamily: pyruvate dehydrogenase (lipoamide) alpha chain; thiamin pyrophosphate-di

C/Keywords: mitochondrion; oxidoreductase

## Query Match

Best Local Similarity 19.7%; Score 39; DB 2; Length 64;  
Matches 6; Conservative 3; Mismatches 4; Indels 0; Gaps 0;

QY 15 PLWGPALKLMKG 27  
DB 11 PYYRERGVLIMRG 23

Search completed: May 6, 2003, 14:58:40  
Job time: 18 secs

GenCore version 5.1.4 p5 4578  
Copyright (c) 1993 - 2003 CompuGen Ltd.

OM protein - protein search, using sw model

Run on: May 6, 2003, 14:57:29 / Search time 16 Seconds  
(without alignments)  
194.145 Million cell updates/sec

Title: US-09-868-399-1

Perfect score: 198

Sequence: 1 KIONFRVYRSDPLMKGPALKMKGGAIVIQDN 36

Scoring table:

BLOSUM62  
Gapop 10.0, Gapext 0.5

Searched: 328255 seqs, 8628685 residues

Total number of hits satisfying chosen parameters: 125019

Minimum DB seq length: 0

Maximum DB seq length: 100

Post-processing: Minimum Match 0%

Maximum Match 100%

Listing first 45 summaries

Database:

Published Applications AA:\*  
1: /cgn2\_6/ptodata/1/pubppaa/US08\_NEW\_PUB.pep:\*  
2: /cgn2\_6/ptodata/1/pubppaa/PCT\_NEW\_PUB.pep:\*  
3: /cgn2\_6/ptodata/1/pubppaa/US06\_NEW\_PUB.pep:\*  
4: /cgn2\_6/ptodata/1/pubppaa/US07\_NEW\_PUB.pep:\*  
5: /cgn2\_6/ptodata/1/pubppaa/US07\_PUBCOMB.pep:\*  
6: /cgn2\_6/ptodata/1/pubppaa/US07\_PUBCOMB.pep:\*  
7: /cgn2\_6/ptodata/1/pubppaa/US07\_PUBCOMB.pep:\*  
8: /cgn2\_6/ptodata/1/pubppaa/US09\_PUBCOMB.pep:\*  
9: /cgn2\_6/ptodata/1/pubppaa/US09\_NEW\_PUB.pep:\*  
10: /cgn2\_6/ptodata/1/pubppaa/US09\_PUBCOMB.pep:\*  
11: /cgn2\_6/ptodata/1/pubppaa/US10\_NEW\_PUB.pep:\*  
12: /cgn2\_6/ptodata/1/pubppaa/US10\_PUBCOMB.pep:\*  
13: /cgn2\_6/ptodata/1/pubppaa/US60\_NEW\_PUB.pep:\*  
14: /cgn2\_6/ptodata/1/pubppaa/US60\_PUBCOMB.pep:\*

Pred. No. is the number of results predicted by chance to have a score greater than or equal to the score of the result being printed, and is derived by analysis of the total score distribution.

#### SUMMARIES

Result No.	Score	Query Match	Length	ID	Description
1	54	27.3	10	US-09-894-018-161	Sequence 161, App
2	54	27.3	15	US-09-894-018-192	Sequence 192, App
3	49	24.7	9	US-10-106-487-36	Sequence 36, Appl
4	49	24.7	14	US-10-062-710-23	Sequence 23, Appl
5	46.5	23.5	96	US-09-764-868-871	Sequence 871, App
6	46	23.2	38	US-10-082-830-200	Sequence 200, App
7	45.5	23.0	79	US-10-125-540-534	Sequence 534, App
8	45.5	23.0	79	US-09-764-870-534	Sequence 534, App
9	44.5	22.5	93	US-09-764-891-4912	Sequence 4912, App
10	44.5	22.5	93	US-09-764-891-5337	Sequence 5337, App
11	43	21.7	75	US-09-764-877-1243	Sequence 1243, App
12	43	21.7	86	US-09-989-920-243	Sequence 243, App
13	41	20.7	53	US-10-011-585A-230	Sequence 230, App
14	41	20.7	53	US-09-864-761-36417	Sequence 36417, A
15	41	20.7	94	US-09-734-569-120	Sequence 120, App
16	40.5	20.5	77	US-09-864-761-48829	Sequence 48829, A
17	40.5	20.5	86	US-09-796-692-2169	Sequence 2169, App
18	40.5	20.5	86	US-10-040-862-2169	Sequence 2169, App
19	40	20.2	31	US-09-822-540A-2	Sequence 2, Appl

20	40	20.2	31	10	US-09-938-700-2	Sequence 2, Appl
21	40	20.2	34	10	US-09-938-700-6	Sequence 6, Appl
22	40	20.2	38	9	US-10-185-050-34	Sequence 34, Appl
23	40	20.2	39	10	US-09-925-297-756	Sequence 756, App
24	40	20.2	60	10	US-09-864-761-49088	Sequence 49088, A
25	40	20.2	63	10	US-09-822-540A-1	Sequence 1, Appl
26	40	20.2	74	10	US-09-925-300-1663	Sequence 1663, App
27	40	20.2	91	9	US-10-001-857-198	Sequence 198, App
28	40	20.2	93	9	US-10-091-504-760	Sequence 760, App
29	40	20.2	93	10	US-09-764-869-760	Sequence 760, App
30	39.5	19.9	57	10	US-09-879-957-213	Sequence 213, App
31	39.5	19.9	63	9	US-09-925-664-34	Sequence 34, Appl
32	39.5	19.9	70	9	US-10-007-280A-210	Sequence 210, App
33	39	19.7	15	10	US-09-894-018-207	Sequence 207, App
34	39	19.7	49	10	US-09-864-761-37881	Sequence 37881, A
35	39	19.7	51	9	US-10-001-835-202	Sequence 202, App
36	39	19.7	66	9	US-09-796-692-2165	Sequence 2165, App
37	39	19.7	66	9	US-10-040-862-2165	Sequence 2165, App
38	39	19.7	67	10	US-09-864-761-33604	Sequence 33604, A
39	39	19.7	81	10	US-09-864-761-47481	Sequence 47481, A
40	39	19.7	88	10	US-09-864-761-46269	Sequence 46269, A
41	38.5	19.4	38	9	US-10-185-050-18	Sequence 18, Appl
42	38.5	19.4	50	10	US-09-864-761-38860	Sequence 38860, A
43	38	19.2	47	9	US-09-809-391-435	Sequence 435, App
44	38	19.2	56	9	US-09-764-891-3292	Sequence 3292, App
45	38	19.2	56	10	US-09-908-711-116	Sequence 116, App

#### ALIGNMENTS

RESULT 1  
US-09-894-018-161  
Sequence 161, Application US/09894018  
Patent No. US20020119127A1  
GENERAL INFORMATION:  
APPLICANT: EPIMUNE, Inc.  
APPLICANT: Sette, Alessandro  
APPLICANT: Chestnut, Robert  
APPLICANT: Livingston, Brian  
APPLICANT: Baker, Denise  
APPLICANT: Newman, Mark  
APPLICANT: Brown, David  
TITLE OF INVENTION: METHODS AND SYSTEM FOR OPTIMIZING  
TITLE OF INVENTION: MINIGENES AND PEPTIDES THEREBY  
FILE REFERENCE: 39963-20033.00  
CURRENT APPLICATION NUMBER: US/09/894,018  
CURRENT FILING DATE: 2001-06-27  
PRIOR APPLICATION NUMBER: PCT/US00/35568  
PRIOR FILING DATE: 2000-12-28  
PRIOR APPLICATION NUMBER: US 60/173,390  
PRIOR FILING DATE: 1999-12-28  
PRIOR APPLICATION NUMBER: US 60/284,221  
PRIOR FILING DATE: 2001-04-16  
NUMBER OF SEQ ID NOS: 368  
SOFTWARE: FastSeq for Windows Version 4.0  
SEQ ID NO 161  
LENGTH: 10  
TYPE: PRT  
ORGANISM: Transgenic mouse  
US-09-894-018-161  
Query Match 27.3%, Score 54, DB 10, Length 10;  
Best Local Similarity 100.0%, Pred. No. 0.23;  
Matches 10; Conservative 0; Mismatches 0; Indels 0; Gaps 0;  
OY 1 KIONFRVYR 10  
|||  
Db 1 KIONFRVYR 10  
RESULT 2  
US-09-894-018-192

```
Sequence 192, Application US/09894018
Patent No. US20020119127A1
GENERAL INFORMATION:
APPLICANT: EPIMUNE, Inc.
APPLICANT: Sette, Alessandro
APPLICANT: Chesnut, Robert
APPLICANT: Livingston, Brian
APPLICANT: Baker, Denise
APPLICANT: Newman, Mark
APPLICANT: Brown, David
TITLE OF INVENTION: METHODS AND SYSTEM FOR OPTIMIZING
TITLE OF INVENTION: ANTIGENS AND PEPTIDES THEREBY
FILE REFERENCE: 39963-2003.00
CURRENT APPLICATION NUMBER: US/09/894,018
PRIOR FILING DATE: 2001-06-27
PRIOR APPLICATION NUMBER: PCT/US00/35568
PRIOR FILING DATE: 2000-12-28
PRIOR APPLICATION NUMBER: US 60/173,390
PRIOR FILING DATE: 1999-12-28
PRIOR APPLICATION NUMBER: US 60/284,221
NUMBER OF SEQ ID NOS: 368
SOFTWARE: FastSeq for Windows Version 4.0
SEQ ID NO 192
LENGTH: 15
TYPE: PRT
ORGANISM: Transgenic mouse
US-09-894-018-192
```

```
Query Match
Best Local Similarity 27.3%; Score 54; DB 10; Length 15;
Matches 10; Conservative 0; Mismatches 0; Indels 0; Gaps 0;
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```
OY 1 KIONRYYR 10
Db 6 KIONRYYR 15
```

```
RESULT 3
US-10-106-487-36
Sequence 36, Application US/10106487
Patent No. US20020164721A1
GENERAL INFORMATION:
APPLICANT: FIRAT, HOSEYIN
APPLICANT: LEMONNIER, FRANCOIS
APPLICANT: LANGLADE-DEMOYEN, PIERRE
APPLICANT: MICHEL, MARIE-LOUISE
TITLE OF INVENTION: DESIGN OF A POLYPEPTIC CONSTRUCT FOR THE INDUCTION
TITLE OF INVENTION: OR
TITLE OF INVENTION: HLA-A2.1 RESTRICTED HIV 1 SPECIFIC CTL RESPONSES USING
FILE REFERENCE: 03495, 0196 SEQUENCE LISTING
CURRENT APPLICATION NUMBER: US/10/106,487
PRIOR FILING DATE: 2002-03-27
PRIOR APPLICATION NUMBER: 09/675,673
PRIOR FILING DATE: 2000-09-29
PRIOR APPLICATION NUMBER: 60/158,356
NUMBER OF SEQ ID NOS: 41
SOFTWARE: PatentIn Ver. 2.1
SEQ ID NO 36
LENGTH: 9
TYPE: PRT
ORGANISM: Human immunodeficiency virus type 1
FEATURE:
NAME/KEY: VARIANT
LOCATION: (9)
OTHER INFORMATION: L9V MUTANT EPILOPE
US-10-106-487-36
```

```
Query Match
Best Local Similarity 24.7%; Score 49; DB 9; Length 9;
Matches 9; Conservative 0; Mismatches 0; Indels 0; Gaps 0;
```

```
OY 23 LMKGEAV 31
Db 1 LMKGEAV 9
```

```
RESULT 4
US-10-062-710-23
Sequence 23, Application US/10062710
Publication No. US20030049253A1
GENERAL INFORMATION:
APPLICANT: Li, Frank O.
APPLICANT: Chu, Yong-Liang
APPLICANT: Qiu, Jian-Tai
TITLE OF INVENTION: Polymeric Conjugates for Delivery of
TITLE OF INVENTION: MHC-Recognized Epitopes
TITLE OF INVENTION: Via Peptide Vaccines
FILE REFERENCE: 3781-001-27
CURRENT APPLICATION NUMBER: US/10/062,710
PRIOR FILING DATE: 2002-02-05
PRIOR APPLICATION NUMBER: US 60/310,498
NUMBER OF SEQ ID NOS: 232
SOFTWARE: FastSeq for Windows Version 4.0
SEQ ID NO 23
LENGTH: 14
TYPE: PRT
ORGANISM: Artificial Sequence
FEATURE:
OTHER INFORMATION: HIV Helper-T Cell Epitopes
US-10-062-710-23
```

```
Query Match
Best Local Similarity 24.7%; Score 49; DB 9; Length 14;
Matches 9; Conservative 0; Mismatches 0; Indels 0; Gaps 0;
```

```
OY 1 KIONRYYR 9
Db 6 KIONRYYR 14
```

```
RESULT 5
US-09-764-868-871
Sequence 871, Application US/09764868
Patent No. US20020168711A1
GENERAL INFORMATION:
APPLICANT: Rosen et al.
TITLE OF INVENTION: Nucleic Acids, Proteins, and Antibodies
FILE REFERENCE: PT232
CURRENT APPLICATION NUMBER: US/09/764,868
PRIOR FILING DATE: 2001-01-17
Prior application data removed - refer to PALM or file wrapper
SOFTWARE: PatentIn Ver. 2.0
SEQ ID NO 871
LENGTH: 96
TYPE: PRT
ORGANISM: Homo sapiens
US-09-764-868-871
```

```
Query Match
Best Local Similarity 23.5%; Score 46.5; DB 9; Length 96;
Matches 13; Conservative 5; Mismatches 11; Indels 9; Gaps 2;
```

```
OY 4 NFRVYRDSRDLKGPAPK-----LW-KGEAV 32
Db 59 NFRVYRDSRDLKGPAPK-----LW-KGEAV 96
```

```
RESULT 6
US-10-082-830-200
Sequence 200, Application US/10082830
Publication No. US2003007604A1
```

```

; APPLICANT: Rosen et al.
; TITLE OF INVENTION: Nucleic Acids, Proteins, and Antibodies
; FILE REFERENCE: PC006
; CURRENT APPLICATION NUMBER: US/09/764,891
; CURRENT FILING DATE: 2001-01-17

```

;; Prior application data removed - consult PALM or file wrapper  
;; NUMBER OF SEQ ID NOS: 10231  
;; SOFTWARE: Patentln Ver. 2.0  
;; SEQ ID NO 5337  
;; LENGTH: 93  
;; TYPE: PRT  
;; ORGANISM: Homo sapiens  
US-09-764-891-5337

Query Match 22.5%; Score 44.5; DB 9; Length 93;  
Best Local Similarity 33.3%; Pred. No. 52;  
Matches 10; Conservative 3; Mismatches 12; Indels 5; Gaps 1;

QY 4 NFRVYRDSRDPIMKGPALKMGKGAIVL 33  
DB 60 NFCIFSRDGVSPCMWG-----NSRTDPLVI 84

RESULT 11  
US-09-764-877-1243  
; Sequence 1243, Application US/09764877  
; Patent No. US20020147140A1  
; GENERAL INFORMATION:  
; APPLICANT: Rosen et al.  
; TITLE OF INVENTION: Nucleic Acids, Proteins, and Antibodies  
; FILE REFERENCE: PC005  
; CURRENT APPLICATION NUMBER: US/09/764,877  
; CURRENT FILING DATE: 2001-01-17  
; Prior application data removed - refer to PALM or file wrapper  
; NUMBER OF SEQ ID NOS: 4031  
; SOFTWARE: Patentln Ver. 2.0  
; SEQ ID NO 1243  
; LENGTH: 75  
; TYPE: PRT  
; ORGANISM: Homo sapiens  
; FEATURE:  
; NAME/KEY: SITE  
; LOCATION: (56)  
; OTHER INFORMATION: Xaa equals any of the naturally occurring L-amino acids  
; NAME/KEY: SITE  
; LOCATION: (75)  
; OTHER INFORMATION: Xaa equals any of the naturally occurring L-amino acids  
US-09-764-877-1243

Query Match 21.7%; Score 43; DB 10; Length 75;  
Best Local Similarity 46.2%; Pred. No. 67;  
Matches \*6; Conservative 4; Mismatches 3; Indels 0; Gaps 0;

QY 8 YYRDSRDPIMKGP 20  
DB 23 YHENTRALIMKGP 35

RESULT 12  
US-09-989-920-243  
; Sequence 243, Application US/09989920  
; Patent No. US20020172957A1  
; GENERAL INFORMATION:  
; APPLICANT: Macina, Roberto  
; APPLICANT: Recipon, Hervé  
; APPLICANT: Chen, Sei-Yu  
; APPLICANT: Sun, Yongming  
; APPLICANT: Liu, Chenghua  
; TITLE OF INVENTION: Compositions and Methods Relating to Lung Specific Genes and Pro  
; FILE REFERENCE: DEX-0291  
; CURRENT APPLICATION NUMBER: US/09/989,920  
; CURRENT FILING DATE: 2001-11-21  
; PRIOR APPLICATION NUMBER: 60/252,500  
; PRIOR FILING DATE: 2000-11-22  
; NUMBER OF SEQ ID NOS: 284  
; SOFTWARE: Patentln version 3.1  
; SEQ ID NO 243  
; LENGTH: 86

;; TYPE: PRT  
;; ORGANISM: Homo sapien  
US-09-989-920-243

Query Match 21.7%; Score 43; DB 9; Length 86;  
Best Local Similarity 50.0%; Pred. No. 77;  
Matches 8; Conservative 1; Mismatches 7; Indels 0; Gaps 0;

QY 4 NFRVYRDSRDPIMKGP 19  
DB 49 NCVFGRDGVSPCMWG 64

RESULT 13  
US-10-011-585A-230  
; Sequence 230, Application US/10011585A  
; Publication No. US2003003986A1  
; GENERAL INFORMATION:  
; APPLICANT: Sun, Yongming  
; APPLICANT: Recipon, Hervé  
; APPLICANT: Chen, Sei-Yu  
; APPLICANT: Liu, Chenghua  
; TITLE OF INVENTION: Genes and Proteins  
; FILE REFERENCE: DEX-0261  
; CURRENT APPLICATION NUMBER: US/10/011,585A  
; CURRENT FILING DATE: 2002-03-14  
; PRIOR APPLICATION NUMBER: 60/245,740  
; PRIOR FILING DATE: 2000-11-03  
; NUMBER OF SEQ ID NOS: 245  
; SOFTWARE: Patentln Ver. 2.1  
; SEQ ID NO 230  
; LENGTH: 38  
; TYPE: PRT  
; ORGANISM: Homo sapiens  
; FEATURE:  
; NAME/KEY: UNSURE  
; LOCATION: (27)  
; OTHER INFORMATION: any amino acid  
US-10-011-585A-230

Query Match 20.7%; Score 41; DB 9; Length 38;  
Best Local Similarity 41.9%; Pred. No. 60;  
Matches 13; Conservative 6; Mismatches 6; Indels 6; Gaps 2;

QY 10 RDSRDPIMKGPALKL---WKGAVIODN 36  
DB 9 RDSRGLKGPGRHMSOXOKGR--VLEDD 37

RESULT 14  
US-09-864-761-36417  
; Sequence 36417, Application US/09864761  
; Patent No. US20020048763A1  
; GENERAL INFORMATION:  
; APPLICANT: Penn, Sharon G.  
; APPLICANT: Rank, David R.  
; APPLICANT: Hanzel, David K.  
; APPLICANT: Chen, Wensheng  
; TITLE OF INVENTION: HUMAN GENOME-DERIVED SINGLE EXON NUCLEIC ACID PROBES USEFUL FOR  
; FILE REFERENCE: Aecmca-X-1  
; CURRENT APPLICATION NUMBER: US/09/864,761  
; CURRENT FILING DATE: 2001-05-23  
; PRIOR APPLICATION NUMBER: US 60/180,312  
; PRIOR FILING DATE: 2000-02-04  
; PRIOR APPLICATION NUMBER: US 60/207,456  
; PRIOR FILING DATE: 2000-05-26  
; PRIOR APPLICATION NUMBER: US 09/632,366  
; PRIOR FILING DATE: 2000-08-03  
; PRIOR APPLICATION NUMBER: GB 24263.6  
; PRIOR FILING DATE: 2000-10-04  
; PRIOR APPLICATION NUMBER: US 60/236,359



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; PRIOR FILING DATE: 2000-09-27
; PRIOR APPLICATION NUMBER: PCT/US01/00666
; PRIOR FILING DATE: 2001-01-30
; PRIOR APPLICATION NUMBER: PCT/US01/00667
; PRIOR FILING DATE: 2001-01-30
; PRIOR APPLICATION NUMBER: PCT/US01/00664
; PRIOR FILING DATE: 2001-01-30
; PRIOR APPLICATION NUMBER: PCT/US01/00669
; PRIOR FILING DATE: 2001-01-30
; PRIOR APPLICATION NUMBER: PCT/US01/00665
; PRIOR FILING DATE: 2001-01-30
; PRIOR APPLICATION NUMBER: PCT/US01/00668
; PRIOR FILING DATE: 2001-01-30
; PRIOR APPLICATION NUMBER: PCT/US01/00663
; PRIOR FILING DATE: 2001-01-30
; PRIOR APPLICATION NUMBER: PCT/US01/00662
; PRIOR FILING DATE: 2001-01-30
; PRIOR APPLICATION NUMBER: PCT/US01/00661
; PRIOR FILING DATE: 2001-01-30
; PRIOR APPLICATION NUMBER: PCT/US01/00670
; PRIOR FILING DATE: 2001-01-30
; PRIOR APPLICATION NUMBER: US 60/234,687
; PRIOR FILING DATE: 2000-09-21
; PRIOR APPLICATION NUMBER: US 09/608,408
; PRIOR FILING DATE: 2000-06-30
; PRIOR APPLICATION NUMBER: US 09/774,203
; PRIOR FILING DATE: 2001-01-29
; NUMBER OF SEQ ID NOS: 49117
; SOFTWARE: Annomax Sequence Listing Engine vers. 1.1
; SEQ ID NO 36417
; LENGTH: 53
; TYPE: PRT
; ORGANISM: Homo sapiens
; FEATURE:
; OTHER INFORMATION: MAP TO AC007322.2
; OTHER INFORMATION: EXPRESSED IN BONE MARROW, SIGNAL = 8.7
; OTHER INFORMATION: EXPRESSED IN BRAIN, SIGNAL = 7.6
; OTHER INFORMATION: EXPRESSED IN PLACENTA, SIGNAL = 5.6
; OTHER INFORMATION: EXPRESSED IN HBL100, SIGNAL = 5.7
; OTHER INFORMATION: EXPRESSED IN HELA, SIGNAL = 4
; OTHER INFORMATION: EXPRESSED IN HEART, SIGNAL = 4.2
; OTHER INFORMATION: EXPRESSED IN FETAL LIVER, SIGNAL = 7
; OTHER INFORMATION: EXPRESSED IN BT474, SIGNAL = 6.5
; OTHER INFORMATION: EXPRESSED IN ADULT LIVER, SIGNAL = 8.7
; OTHER INFORMATION: EXPRESSED IN LUNG, SIGNAL = 7.6
; OTHER INFORMATION: EST_HUMAN HIT: AA196978.1, EVALU6 3.30e-01
US-09-864-761-36417

Query Match 20.7%; Score 41; DB 10; Length 53;
Best Local Similarity 58.3%; Pred. No. 86;
Matches 7; Conservative 2; Mismatches 3; Indels 0; Gaps 0;

QY 25 WKGCAGVVDN 36
Db 1 WKGSAAVLIADH 12

RESULT 15
US-09-734-569-120
; Sequence 120, Application US/09734569
; Patent No. US20020064816A1
; GENERAL INFORMATION:
; APPLICANT: Lerchl, Jens
; APPLICANT: Renz, Andreas
; APPLICANT: Enhardt, Thomas
; APPLICANT: Reindl, Andreas
; APPLICANT: Cifrus, Petra
; APPLICANT: Bischoff, Friedrich
; APPLICANT: Frank, Markus
; APPLICANT: Freund, Annette
; APPLICANT: Duwenig, Elke
; APPLICANT: Schmidt, Ralf-Michael
; APPLICANT: Reekl, Ralf
```

```
; TITLE OF INVENTION: Moss genes from Physcomitrella patens encoding proteins involved
; in the synthesis of carbohydrates
; FILE REFERENCE: BASF-NAE-1332-99-US
; CURRENT APPLICATION NUMBER: US/09/734,569
; CURRENT FILING DATE: 2001-05-24
; PRIOR APPLICATION NUMBER: US 60/171,101
; PRIOR FILING DATE: 1999-12-16
; NUMBER OF SEQ ID NOS: 181
; SOFTWARE: PatentIn Ver. 2.1/WordPerfect 6.1
; SEQ ID NO 120
; LENGTH: 94
; TYPE: PRT
; ORGANISM: Physcomitrella patens
US-09-734-569-120

Query Match 20.7%; Score 41; DB 10; Length 94;
Best Local Similarity 40.6%; Pred. No. 1.6e+02;
Matches 13; Conservative 4; Mismatches 11; Indels 4; Gaps 1;

QY 7 VYRDSRDP---LWKGPALKLWKGAGVTV 34
Db 35 VKYGSMDAFKQILAKGSAKSLFKGAGNIIK 66

Search completed: May 6, 2003, 14:59:23
Job time : 17 secs
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GenCore version 5.1.4 p5 4578  
Copyright (c) 1993 - 2003 Compugen Ltd.

OM protein - protein search, using sw model

Run on: May 6, 2003, 14:56:49 / Search time 15 Seconds  
(without alignments)  
70.615 Million cell updates/sec

Title: US-09-868-399-1  
Perfect score: 198  
Sequence: 1 KIGNFRVYRDSRDLPMKGPALKLMKGEGAVIQDN 36

Scoring table: BLOSUM62  
Gapop 10.0, Gapext 0.5

Searched: 262574 seqs, 29422922 residues

Total number of hits satisfying chosen parameters: 196705

Minimum DB seq length: 0  
Maximum DB seq length: 100

Post-processing: Minimum Match 0%  
Maximum Match 100%

Listing first 45 summaries

Database :

1: Issued\_Patents\_AA.\*  
2: /cgn2\_6/ptodata/1/1aa/5A.COMB.pep.\*  
3: /cgn2\_6/ptodata/1/1aa/5B.COMB.pep.\*  
4: /cgn2\_6/ptodata/1/1aa/6A.COMB.pep.\*  
5: /cgn2\_6/ptodata/1/1aa/PTUG.COMB.pep.\*  
6: /cgn2\_6/ptodata/1/1aa/backfile1.pep.\*

Pred. No. is the number of results predicted by chance to have a score greater than or equal to the score of the result being printed, and is derived by analysis of the total score distribution.

## SUMMARIES

Result No.	Score	Query Match	Length	ID	Description
1	79	39.9	15	4	US-09-009-953-221
2	58	29.3	43	2	US-08-468-161-67
3	58	29.3	43	2	US-09-273-685-67
4	58	29.3	43	5	PCT-US95-11934-67
5	54	27.3	10	3	US-08-159-339A-550
6	54	27.3	15	4	US-09-009-953-220
7	53	26.8	9	2	US-08-986-234-71
8	53	26.8	10	4	US-08-197-484-74
9	53	26.8	10	5	PCT-US95-02121-74
10	49	24.7	9	2	US-08-986-234-72
11	49	24.7	9	4	US-08-197-484-75
12	49	24.7	9	5	PCT-US95-02121-75
13	49	24.7	11	3	US-08-159-339A-1140
14	48	24.2	15	4	US-09-255-502-6
15	46	23.2	43	3	US-08-468-161-51
16	46	23.2	43	3	US-09-273-685-51
17	46	23.2	43	5	PCT-US95-11934-51
18	44	22.2	30	4	US-09-315-304B-1521
19	44	22.2	67	3	US-08-475-668A-210
20	44	22.2	67	3	US-08-485-551A-210
21	44	22.2	77	3	US-08-475-668A-211
22	44	22.2	77	3	US-08-485-551A-211
23	43	21.7	34	1	US-08-460-602A-26
24	43	21.7	34	1	US-08-463-966A-26
25	43	21.7	34	1	US-08-465-217A-26
26	43	21.7	34	2	US-08-464-329A-26
27	43	21.7	34	2	US-08-462-507A-26

28	43	21.7	34	2	US-08-462-507A-26	Sequence 26, App1
29	43	21.7	34	2	US-08-467-881A-26	Sequence 26, App1
30	43	21.7	34	4	US-08-750-624-4	Sequence 4, App1
31	43	21.7	37	1	US-08-257-528B-39	Sequence 39, App1
32	43	21.7	37	1	US-08-460-602A-39	Sequence 39, App1
33	43	21.7	37	1	US-08-463-966A-39	Sequence 39, App1
34	43	21.7	37	1	US-08-465-217A-39	Sequence 39, App1
35	43	21.7	37	2	US-08-464-329A-39	Sequence 39, App1
36	43	21.7	37	2	US-08-462-507A-39	Sequence 39, App1
37	43	21.7	37	2	US-08-467-881A-39	Sequence 39, App1
38	43	21.7	40	1	US-08-426-819A-19	Sequence 19, App1
39	41.5	21.0	66	4	US-09-082-593-6	Sequence 6, App1
40	41	20.7	44	1	US-08-257-528B-95	Sequence 95, App1
41	41	20.7	44	1	US-08-460-602A-95	Sequence 95, App1
42	41	20.7	44	1	US-08-463-966A-95	Sequence 95, App1
43	41	20.7	44	1	US-08-465-217A-95	Sequence 95, App1
44	41	20.7	44	2	US-08-464-329A-95	Sequence 95, App1
45	41	20.7	44	2	US-08-462-507A-95	Sequence 95, App1

## ALIGNMENTS

RESULT 1  
US-09-009-953-221  
Sequence 221, Application US/09009953  
Patent No. 6413517  
GENERAL INFORMATION:  
APPLICANT: Sette, Alessandro  
TITLE OF INVENTION: Identification of Broadly Reactive DR Restricted Epitopes  
NUMBER OF SEQUENCES: 274  
CORRESPONDENCE ADDRESS:  
ADDRESSEE: Townsend and Townsend and Crew LLP  
STREET: Two Embarcadero Center, Eighth Floor  
CITY: San Francisco  
STATE: CA  
COUNTRY: USA  
ZIP: 94111-3834  
COMPUTER READABLE FORM:  
MEDIUM TYPE: Diskette  
COMPUTER: IBM Compatible  
OPERATING SYSTEM: DOS  
SOFTWARE: FastSeq for Windows Version 2.0  
CURRENT APPLICATION DATA:  
APPLICATION NUMBER: US/09/009,953  
FILING DATE: 21-Jan-1998  
CLASSIFICATION: <Unknown>  
PRIOR APPLICATION DATA:  
APPLICATION NUMBER: US 60/036,713  
FILING DATE: 23-JAN-1997  
APPLICATION NUMBER: US 60/037,432  
FILING DATE: 07-FEB-1997  
ATTORNEY/AGENT INFORMATION:  
NAME: Weber, Ellen Lauver  
REGISTRATION NUMBER: 32,762  
REFERENCE/DOCKET NUMBER: 018623-011520US  
TELECOMMUNICATION INFORMATION:  
TELEPHONE: 415-576-0200  
TELEFAX: 415-576-0300  
TELEX: <Unknown>  
INFORMATION FOR SEQ ID NO: 221:  
SEQUENCE CHARACTERISTICS:  
LENGTH: 15 amino acids  
TYPE: amino acid  
STRANDEDNESS: single  
TOPOLOGY: linear  
MOLECULE TYPE: peptide  
SEQUENCE DESCRIPTION: SEQ ID NO: 221:  
US-09-009-953-221  
Query Match 39.9%; Score 79; DB 4; Length 15;  
Best Local Similarity 100.0%; Pred. No. 1.9e-05;

Matches 15; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

QY 22 KILWKGAVIION 36

Db 1 KILWKGAVIION 15

## RESULT 2

US-08-488-161-67

; Sequence 67, Application US/08488161  
; Patent No. 5885577

; GENERAL INFORMATION:

; APPLICANT: Alvarez, Vernon L.

; TITLE OF INVENTION: Antigen Binding Peptides (Abtides) From

; NUMBER OF SEQUENCES: 103

; CORRESPONDENCE ADDRESS:

; ADDRESSEE: Pennie & Edmonds

; STREET: 1155 Avenue of the Americas

; CITY: New York

; STATE: New York

; COUNTRY: USA

; ZIP: 10036

; COMPUTER READABLE FORM:

; MEDIUM TYPE: Floppy disk

; OPERATING SYSTEM: IBM PC compatible

; SOFTWARE: Patentin Release #1.0, Version #1.30

; CURRENT APPLICATION DATA:

; APPLICATION NUMBER: US/08/488,161

; FILING DATE: 07-JUN-1995

; CLASSIFICATION: 436

; ATTORNEY/AGENT INFORMATION:

; NAME: Mistrock, S. Leslie

; REGISTRATION NUMBER: 18,872

; REFERENCE/DOCKET NUMBER: 1101-176

; TELECOMMUNICATION INFORMATION:

; TELEPHONE: (212) 790-9090

; TELEFAX: (212) 869-9741/8864

; TELEX: 66141 PENNIE

; INFORMATION FOR SEQ ID NO: 67:

; SEQUENCE CHARACTERISTICS:

; LENGTH: 43 amino acids

; TYPE: amino acid

; STRANDEDNESS: single

; TOPOLOGY: linear

; MOLECULE TYPE: peptide

; US-08-488-161-67

Query Match 29.3%; Score 58; DB 2; Length 43;

Best Local Similarity 66.7%; Pred. No. 0.083;

Matches 10; Conservative 1; Mismatches 4; Indels 0; Gaps 0;

QY 15 PLWKGPAKILWKGEG 29

Db 9 PPMGSPAGILWGGCG 23

RESULT 3

US-09-273-685-67

; Sequence 67, Application US/09273685

; Patent No. 6015561

; GENERAL INFORMATION:

; APPLICANT: Alvarez, Vernon L.

; TITLE OF INVENTION: Antigen Binding Peptides (Abtides) From

; NUMBER OF SEQUENCES: 103

; CORRESPONDENCE ADDRESS:

; ADDRESSEE: Pennie & Edmonds

; STREET: 1155 Avenue of the Americas

; CITY: New York

; STATE: New York

; COUNTRY: USA

ZIP: 10036

; COMPUTER READABLE FORM:

; MEDIUM TYPE: Floppy disk

; OPERATING SYSTEM: IBM PC compatible

; SOFTWARE: Patentin Release #1.0, Version #1.30

; CURRENT APPLICATION DATA:

; APPLICATION NUMBER: US/09/273,685

; FILING DATE:

; CLASSIFICATION:

; PRIOR APPLICATION DATA:

; APPLICATION NUMBER: 08/488,161

; FILING DATE:

; ATTORNEY/AGENT INFORMATION:

; NAME: Mistrock, S. Leslie

; REGISTRATION NUMBER: 18,872

; REFERENCE/DOCKET NUMBER: 1101-176

; TELECOMMUNICATION INFORMATION:

; TELEPHONE: (212) 790-9090

; TELEFAX: (212) 869-9741/8864

; TELEX: 66141 PENNIE

; INFORMATION FOR SEQ ID NO: 67:

; SEQUENCE CHARACTERISTICS:

; LENGTH: 43 amino acids

; TYPE: amino acid

; STRANDEDNESS: single

; TOPOLOGY: linear

; MOLECULE TYPE: peptide

; US-09-273-685-67

Query Match 29.3%; Score 58; DB 3; Length 43;

Best Local Similarity 66.7%; Pred. No. 0.083;

Matches 10; Conservative 1; Mismatches 4; Indels 0; Gaps 0;

QY 15 PLWKGPAKILWKGEG 29

Db 9 PPMGSPAGILWGGCG 23

RESULT 4

PCT-US95-11934-67

; Sequence 67, Application PC/TUS9511934

; GENERAL INFORMATION:

; APPLICANT: Cytogen Corporation

; TITLE OF INVENTION: Antigen Binding Peptides (Abtides) From

; NUMBER OF SEQUENCES: 103

; CORRESPONDENCE ADDRESS:

; ADDRESSEE: Pennie & Edmonds

; STREET: 1155 Avenue of the Americas

; CITY: New York

; STATE: New York

; COUNTRY: USA

; ZIP: 10036

; COMPUTER READABLE FORM:

; MEDIUM TYPE: Floppy disk

; OPERATING SYSTEM: IBM PC compatible

; SOFTWARE: Patentin Release #1.0, Version #1.30

; CURRENT APPLICATION DATA:

; APPLICATION NUMBER: PCT/US95/11934

; FILING DATE: 20-SEP-1995

; CLASSIFICATION:

; ATTORNEY/AGENT INFORMATION:

; NAME: Mistrock, S. Leslie

; REGISTRATION NUMBER: 18,872

; REFERENCE/DOCKET NUMBER: 1101-196-228

; TELECOMMUNICATION INFORMATION:

; TELEPHONE: (212) 790-9090

; TELEFAX: (212) 869-9741/8864

; TELEX: 66141 PENNIE

; INFORMATION FOR SEQ ID NO: 67:

; SEQUENCE CHARACTERISTICS:

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; LENGTH: 43 amino acids
; TYPE: amino acid
; STRANDEDNESS: single
; TOPOLOGY: linear
; MOLECULE TYPE: peptide
PCT-US95-11934-67

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Query Match	29.3%;	Score 58;	DB 5;	Length 43;
Best Local Similarity	66.7%;	Pred. No. 0.083;	.	
Matches 10;	Conservative	1;	Mismatches	4;
			Indels	

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QY      15 PLWKGPAKLWKGEQ 29
          |||||:|
Db      9  PWSGPAGLWQGC 23
```

RESULT 5  
US-08-159-339A-550  
; Sequence 550, Application US/08159339A

1 GENERAL INFORMATION:  
2 APPLICANT: Kubo, Ralph T.  
3 APPLICANT: Grey, Howard M.  
4 APPLICANT: Sette, Alessandro  
5 APPLICANT: Celis, Esteban  
6 TITLE OF INVENTION: HLA Binding peptides and Their  
7 TITLE OF INVENTION: Uses  
8 NUMBER OF SEQUENCES: 1254  
9 CORRESPONDENCE ADDRESS:  
0

```

; COUNTRY: USA
; ZIP: 94111-3834
; COMPUTER READABLE FORM:
;

```

COMPUTER: IBM Compatible  
OPERATING SYSTEM: DOS  
SOFTWARE: FastEED for Windows Vers 1.0  
CURRENT APPLICATION DATA:  
APPLICATION NUMBER: US/08/159,339A  
FILING DATE: 29-NOV-1993  
CLASSIFICATION: 424  
PRIOR APPLICATION DATA:

1  
2  
3 APPLICATION NUMBER: US 07/926,566  
4  
5 FILING DATE: 07-AUG-1992  
6  
7 APPLICATION NUMBER: US 08/027,746  
8  
9 FILING DATE: 05-MAR-1993  
10  
11 APPLICATION NUMBER: US 08/103,396  
12  
13 FILING DATE: 06-AUG-1993  
14  
15 ATTORNEY/AGENT INFORMATION:  
16

```

NAME: Weber, Ellen Luavei
REGISTRATION NUMBER: 32,762
REFERENCE/DOCKET NUMBER: 018623-005030US
TELECOMMUNICATION INFORMATION:
TELEPHONE: (415) 576-0200
TELEFAX: (415) 576-0300
TELEX:
INFORMATION FOR SEQ ID NO: 550:
SEQUENCE CHARACTERISTICS:
LENGTH: 10 amino acids
TYPE: amino acid
STRANDEDNESS: single
TOPOLOGY: linear
MOLECULE TYPE: peptide
OS-08-159-339A-550

```

Query Match	27.3%;	Score 54;	DB 3;	Length 10;
Best Local Similarity	100.0%;	Pred. No. 0.058;		
Matches 10;	Conservative 0;	Mismatches 0;	Indels	

QY 1 KI QNFRVYYR 10

Db 1 KIÖNFRVYYR 10

RESULT 6  
US-09-009-953-220  
; Sequence 220, Application US/09009953  
; Patent No. 6413517  
; GENERAL INFORMATION:  
; Application, Status, Information

TITLE OF INVENTION: Identification of Broadly  
 Reactive DR Restricted Epitopes  
 NUMBER OF SEQUENCES: 274  
 CORRESPONDENCE ADDRESS:  
 ADDRESSEE: Townsend and Crew LLP  
 STREET: Two Embarcadero Center, Eighth Floor  
 CITY: San Francisco  
 STATE: CA

```

?      COUNTRY: USA
?      ZIP: 94111-3834
?
?      COMPUTER READABLE FORM:
?      MEDIUM TYPE: Diskette
?      COMPUTER: IBM Compatible
?      OPERATING SYSTEM: DOS
?      SOFTWARE: PASSED for Windows Version 2.0
?
?      CURRENT APPLICATION DATA:
?      APPLICATION NUMBER: US/09/009,953

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CLASSIFICATION: <Unknown>  
PRIOR APPLICATION DATA:  
APPLICATION NUMBER: US 60/036,713  
FILING DATE: 23-JAN-1997  
APPLICATION NUMBER: US 60/037,432  
FILING DATE: 07-FEB-1997  
ATTORNEY/AGENT INFORMATION:  
NAME: Weber, Ellen Lawyer  
REGISTRATION NUMBER: 32,762  
REFERENCE/DOCKET NUMBER: 018623-011520US  
TELECOMMUNICATION INFORMATION:

```

?      TELEFAX: 415-576--0300
?      TELEX : <Unknown>
?      INFORMATION FOR SEQ ID NO: 220:
?          SEQUENCE CHARACTERISTICS:
?              LENGTH: 15 amino acids
?                  TYPE: amino acid
?                      STRANDEDNESS: single
?                          TOPOLOGY: linear
?                              MOLECULE TYPE: peptide
?                                  ?
?                                  ?
?                                  ?
US-09-009-953-220

Query Match           27.3%; Score 54; DB 4; Length 15;
Best Local Similarity 100.0%; Pred. No. 0.094;
Matches    10; Conservative    0; Mismatches    0; Indels    0; Gaps    0

```

Qy	1	KIQNFRVYYR	10
Db	6	KIQNFRVYYR	15

RESULT 7  
US-08-986-234-71  
; Sequence 71, Applicat

```

RESULT 7
US-08-986-234-71
; Sequence 71, Application US/08986234
; Patent No. 5981706
; GENERAL INFORMATION:
; APPLICANT: Wellen, et al.
; TITLE OF INVENTION: Methods for Synthesizing Heat Shock Protein Complexes
; FILE REFERENCE: UNNE-0008-1
; CURRENT APPLICATION NUMBER: US/08/986,234
; CURRENT FILING DATE: 1997-12-05
; NUMBER OF SEQ ID NOS: 114
; SOFTWARE: PatentIn Ver. 2.0

```

SEQ ID NO 71  
LENGTH: 9  
TYPE: PRT  
ORGANISM: Human immunodeficiency virus  
US-08-986-234-71

Query Match 26.8%; Score 53; DB 2; Length 9;  
Best Local Similarity 100.0%; Pred. No. 1.9e+05;  
Matches 9; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

QY 15 LLMKGPATL 23  
Db 1 LLMKGPATL 9

RESULT 8  
US-08-197-484-74  
Sequence 74, Application US/08197484  
Patent No. 6419931  
GENERAL INFORMATION:  
APPLICANT: VITTELO, Maria A.  
APPLICANT: CHESTNUT, Robert W.  
APPLICANT: SETTE, Alessandro D.  
APPLICANT: CELIS, Esteban  
APPLICANT: GRAY, Howard  
TITLE OF INVENTION: COMPOSITIONS AND METHODS FOR ELICITING  
TITLE OF INVENTION: CTL IMMUNITY  
NUMBER OF SEQUENCES: 153  
CORRESPONDENCE ADDRESS:  
ADDRESSEE: Townsend and Townsend Kourie and Crew  
STREET: Steuart Street Tower, One Market Plaza  
CITY: San Francisco  
STATE: California  
COUNTRY: US  
ZIP: 94105-1493  
COMPUTER READABLE FORM:  
MEDIUM TYPE: Floppy disk  
COMPUTER: IBM PC compatible  
OPERATING SYSTEM: PC-DOS/MS-DOS  
SOFTWARE: Patentin Release #1.0, Version #1.25  
CURRENT APPLICATION DATA:  
APPLICATION NUMBER: US/08/197,484  
FILING DATE: 16-FEB-1994  
CLASSIFICATION: 424  
PRIOR APPLICATION DATA:  
APPLICATION NUMBER: US 07/935,811  
FILING DATE: 26-AUG-1992  
PRIOR APPLICATION DATA:  
APPLICATION NUMBER: US 07/874,491  
FILING DATE: 27-APR-1992  
PRIOR APPLICATION DATA:  
APPLICATION NUMBER: US 07/827,682  
FILING DATE: 29-JAN-1992  
PRIOR APPLICATION DATA:  
APPLICATION NUMBER: US 07/749,568  
FILING DATE: 26-AUG-1991  
ATTORNEY/AGENT INFORMATION:  
NAME: Parmelee, Steven W.  
REGISTRATION NUMBER: 31,990  
REFERENCE/DOCKET NUMBER: 14137-26-4  
TELECOMMUNICATION INFORMATION:  
TELEPHONE: (206) 467-9600  
TELEFAX: (206) 467-9600  
INFORMATION FOR SEQ ID NO: 74:  
SEQUENCE CHARACTERISTICS:  
LENGTH: 10 amino acids  
TYPE: amino acid  
STRANDEDNESS: unknown  
TOPOLOGY: unknown  
MOLECULE TYPE: peptide  
US-08-197-484-74

Query Match 26.8%; Score 53; DB 4; Length 10;

Best Local Similarity 100.0%; Pred. No. 0.082;  
Matches 10; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

QY 23 LLMKGEAVV 32  
Db 1 LLMKGEAVV 10

RESULT 9  
PCT-US95-02121-74  
Sequence 74, Application PC/TUS9502121  
GENERAL INFORMATION:  
APPLICANT:  
TITLE OF INVENTION: COMPOSITIONS AND METHODS FOR ELICITING  
TITLE OF INVENTION: CTL IMMUNITY  
NUMBER OF SEQUENCES: 153  
COMPUTER READABLE FORM:  
MEDIUM TYPE: Floppy disk  
COMPUTER: IBM PC compatible  
OPERATING SYSTEM: PC-DOS/MS-DOS  
SOFTWARE: Patentin Release #1.0, Version #1.25  
CURRENT APPLICATION DATA:  
APPLICATION NUMBER: PCT/US95/02121  
FILING DATE: 16-FEB-1995  
CLASSIFICATION:  
PRIOR APPLICATION DATA:  
APPLICATION NUMBER: US 08/197,484  
FILING DATE: 16-FEB-1994  
PRIOR APPLICATION DATA:  
APPLICATION NUMBER: US 07/935,811  
FILING DATE: 26-AUG-1992  
PRIOR APPLICATION DATA:  
APPLICATION NUMBER: US 07/874,491  
FILING DATE: 27-APR-1992  
PRIOR APPLICATION DATA:  
APPLICATION NUMBER: US 07/827,682  
FILING DATE: 29-JAN-1992  
PRIOR APPLICATION DATA:  
APPLICATION NUMBER: US 07/749,568  
FILING DATE: 26-AUG-1991  
ATTORNEY/AGENT INFORMATION:  
NAME: Parmelee, Steven W.  
REGISTRATION NUMBER: 31,990  
REFERENCE/DOCKET NUMBER: 14137-26-4PC  
TELECOMMUNICATION INFORMATION:  
TELEPHONE: (206) 543-5043  
TELEFAX: (206) 467-9600  
INFORMATION FOR SEQ ID NO: 74:  
SEQUENCE CHARACTERISTICS:  
LENGTH: 10 amino acids  
TYPE: amino acid  
STRANDEDNESS: unknown  
TOPOLOGY: unknown  
MOLECULE TYPE: peptide  
PCT-US95-02121-74

Query Match 26.8%; Score 53; DB 5; Length 10;  
Best Local Similarity 100.0%; Pred. No. 0.082;  
Matches 10; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

QY 23 LLMKGEAVV 32  
Db 1 LLMKGEAVV 10

RESULT 10  
US-08-986-234-72  
Sequence 72, Application US/08986234  
Patent No. 5981706  
GENERAL INFORMATION:  
APPLICANT: Wallen, et al.  
TITLE OF INVENTION: Methods for Synthesizing Heat Shock Protein Complexes  
FILE REFERENCE: UNME-0008-1

MOLECULE TYPE: peptide  
US-08-197-484-75  
Query Match 24.7%; Score 49; DB 2; Length 9;  
Best Local Similarity 100.0%; Pred. No. 1.9e+05;  
Matches 9; Conservative 0; Mismatches 0; Indels 0; Gaps 0;  
OY 23 LMKGEAV 31  
DB 1 LMKGEAV 9  
RESULT 11  
US-08-197-484-75  
Sequence 75, Application US/08197484  
Patent No. 6419931  
GENERAL INFORMATION:  
APPLICANT: VITTELLO, Maria A.  
APPLICANT: CHESTNUT, Robert W.  
APPLICANT: SETTE, Alessandro D.  
APPLICANT: CELIS, Estebean  
APPLICANT: GRAY, Howard  
TITLE OF INVENTION: COMPOSITIONS AND METHODS FOR ELICITING  
TITLE OF INVENTION: CTL IMMUNITY  
NUMBER OF SEQUENCES: 153  
CORRESPONDENCE ADDRESS:  
ADDRESSEE: Townsend and Townsend Kourie and Crew  
STREET: Steuart Street Tower, One Market Plaza  
CITY: San Francisco  
STATE: California  
COUNTRY: US  
ZIP: 94105-1493  
COMPUTER READABLE FORM:  
MEDIUM TYPE: Floppy disk  
COMPUTER: IBM PC compatible  
OPERATING SYSTEM: PC-DOS/MS-DOS  
SOFTWARE: Patentin Release #1.0, Version #1.25  
CURRENT APPLICATION DATA:  
APPLICATION NUMBER: US/08/197,484  
FILING DATE: 16-FEB-1994  
CLASSIFICATION: 424  
PRIOR APPLICATION DATA:  
APPLICATION NUMBER: US 07/935,811  
FILING DATE: 26-AUG-1992  
PRIOR APPLICATION DATA:  
APPLICATION NUMBER: US 07/874,491  
FILING DATE: 27-APR-1992  
PRIOR APPLICATION DATA:  
APPLICATION NUMBER: US 07/827,682  
FILING DATE: 29-JAN-1992  
PRIOR APPLICATION DATA:  
APPLICATION NUMBER: US 07/749,568  
FILING DATE: 26-AUG-1991  
ATTORNEY/AGENT INFORMATION:  
NAME: Parmelee, Steven W.  
REGISTRATION NUMBER: 31,990  
REFERENCE/DOCKET NUMBER: 14137-26-4  
TELECOMMUNICATION INFORMATION:  
TELEPHONE: (206) 467-9600  
TELEFAX: (206) 623-6793  
INFORMATION FOR SEQ ID NO: 75:  
SEQUENCE CHARACTERISTICS:  
LENGTH: 9  
TYPE: amino acid  
STRANDEDNESS: unknown  
TOPOLOGY: unknown

MOLECULE TYPE: peptide  
US-08-197-484-75  
Query Match 24.7%; Score 49; DB 4; Length 9;  
Best Local Similarity 100.0%; Pred. No. 1.9e+05;  
Matches 9; Conservative 0; Mismatches 0; Indels 0; Gaps 0;  
OY 23 LMKGEAV 31  
DB 1 LMKGEAV 9  
RESULT 12  
PCT-US95-02121-75  
Sequence 75, Application PC/TUS9502121  
GENERAL INFORMATION:  
APPLICANT:  
TITLE OF INVENTION: COMPOSITIONS AND METHODS FOR ELICITING  
TITLE OF INVENTION: CTL IMMUNITY  
NUMBER OF SEQUENCES: 153  
COMPUTER READABLE FORM:  
MEDIUM TYPE: Floppy disk  
COMPUTER: IBM PC compatible  
OPERATING SYSTEM: PC-DOS/MS-DOS  
SOFTWARE: Patentin Release #1.0, Version #1.25  
CURRENT APPLICATION DATA:  
APPLICATION NUMBER: PCT/US95/02121  
FILING DATE: 16-FEB-1995  
CLASSIFICATION:  
PRIOR APPLICATION DATA:  
APPLICATION NUMBER: US 08/197,484  
FILING DATE: 16-FEB-1994  
PRIOR APPLICATION DATA:  
APPLICATION NUMBER: US 07/935,811  
FILING DATE: 26-AUG-1992  
PRIOR APPLICATION DATA:  
APPLICATION NUMBER: US 07/874,491  
FILING DATE: 27-APR-1992  
PRIOR APPLICATION DATA:  
APPLICATION NUMBER: US 07/827,682  
FILING DATE: 29-JAN-1992  
PRIOR APPLICATION DATA:  
APPLICATION NUMBER: US 07/749,568  
FILING DATE: 26-AUG-1991  
ATTORNEY/AGENT INFORMATION:  
NAME: Parmelee, Steven W.  
REGISTRATION NUMBER: 31,990  
REFERENCE/DOCKET NUMBER: 14137-26-4PC  
TELECOMMUNICATION INFORMATION:  
TELEPHONE: (206) 467-9600  
TELEFAX: (415) 543-5043  
INFORMATION FOR SEQ ID NO: 75:  
SEQUENCE CHARACTERISTICS:  
LENGTH: 9  
TYPE: amino acid  
STRANDEDNESS: unknown  
TOPOLOGY: unknown  
MOLECULE TYPE: peptide  
PCT-US95-02121-75  
Query Match 24.7%; Score 49; DB 5; Length 9;  
Best Local Similarity 100.0%; Pred. No. 1.9e+05;  
Matches 9; Conservative 0; Mismatches 0; Indels 0; Gaps 0;  
OY 23 LMKGEAV 31  
DB 1 LMKGEAV 9  
RESULT 13  
US-08-159-339A-1140  
Sequence 1140, Application US/08159339A  
Patent No. 6037135

GENERAL INFORMATION:  
APPLICANT: Kubo, Ralph T.  
APPLICANT: Grey, Howard M.  
APPLICANT: Sette, Alessandro  
APPLICANT: Celis, Esteban  
TITLE OF INVENTION: HLA Binding peptides and Their  
TITLE OF INVENTION: Uses  
NUMBER OF SEQUENCES: 1254  
CORRESPONDENCE ADDRESS:  
ADDRESSEE: Townsend and Townsend and Crew LLP  
STREET: Two Embarcadero Center, Eighth Floor  
CITY: San Francisco  
STATE: CA  
COUNTRY: USA  
ZIP: 94111-3834  
COMPUTER READABLE FORM:  
MEDIUM TYPE: Diskette  
COMPUTER: IBM Compatible  
OPERATING SYSTEM: DOS  
SOFTWARE: FastSeq for Windows Version 2.0  
CURRENT APPLICATION DATA:  
APPLICATION NUMBER: US/08/159,339A  
FILING DATE: 29-NOV-1993  
CLASSIFICATION: 424  
PRIOR APPLICATION DATA:  
APPLICATION NUMBER: US 07/926,666  
FILING DATE: 07-AUG-1992  
APPLICATION NUMBER: US 08/027,746  
FILING DATE: 05-MAR-1993  
APPLICATION NUMBER: US 08/103,396  
FILING DATE: 06-AUG-1993  
ATTORNEY/AGENT INFORMATION:  
NAME: Weber, Ellen Lauver  
REGISTRATION NUMBER: 32,762  
REFERENCE/DOCKET NUMBER: 018623-005030US  
TELECOMMUNICATION INFORMATION:  
TELEPHONE: (415) 576-0200  
TELEFAX: (415) 576-0300  
TELEX:  
INFORMATION FOR SEQ ID NO: 1140:  
SEQUENCE CHARACTERISTICS:  
LENGTH: 11 amino acids  
TYPE: amino acid  
STRANDEDNESS: single  
TOPOLOGY: linear  
MOLECULE TYPE: peptide  
US-08-159-339A-1140  
Query Match 24.7%; Score 49; DB 3; Length 11;  
Best Local Similarity 100.0%; Pred. No. 0.36;  
Matches 9; Conservative 0; Mismatches 0; Indels 0; Gaps 0;  
QY 1 KIONFRVY 9.  
|||  
Db 3 KIONFRVY 11  
RESULT 14  
US-09-255-502-6  
Sequence 6, Application US/09255502  
Patent No. 6218165  
GENERAL INFORMATION:  
APPLICANT: Estell, David  
APPLICANT: Harding, Fiona  
TITLE OF INVENTION: Mutant Proteins Having Lower Allergenic Responses in  
TITLE OF INVENTION: Humans and Methods for Constructing, Identifying and  
FILE REFERENCE: GC 527-D2  
CURRENT APPLICATION NUMBER: US/09/255,502  
PRIOR FILING DATE: 1999-02-23  
CURRENT APPLICATION NUMBER: 09/060,872  
PRIOR FILING DATE: 1998-04-15  
NUMBER OF SEQ ID NOS: 7

SOFTWARE: PatentIn Ver. 2.0  
SEQ ID NO 6  
LENGTH: 15  
TYPE: PRT  
ORGANISM: Unknown  
FEATURE:  
OTHER INFORMATION: Description of Unknown Organism: Unknown Sequence  
US-09-255-502-6

Query Match 24.2%; Score 48; DB 4; Length 15;  
Best Local Similarity 61.5%; Pred. No. 0.72;  
Matches 8; Conservative 4; Mismatches 1; Indels 0; Gaps 0;

QY 2 IONFRVYRDSRD 14  
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Db 1 IONFRVYRDSRD 13

RESULT 15  
US-08-488-161-51  
Sequence 51, Application US/08488161  
Patent No. 5885577  
GENERAL INFORMATION:  
APPLICANT: Alvarez, Vernon L.  
TITLE OF INVENTION: Antigen Binding Peptides (Abtides) From  
NUMBER OF SEQUENCES: 103  
CORRESPONDENCE ADDRESS:  
ADDRESSEE: Pennie & Edmonds  
STREET: 1155 Avenue of the Americas  
CITY: New York  
STATE: New York  
COUNTRY: USA  
ZIP: 10036  
COMPUTER READABLE FORM:  
MEDIUM TYPE: Floppy disk  
COMPUTER: IBM PC compatible  
OPERATING SYSTEM: PC-DOS/MS-DOS  
SOFTWARE: PatentIn Release #1.0, Version #1.30  
CURRENT APPLICATION DATA:  
APPLICATION NUMBER: US/08/488,161  
FILING DATE: 07-JUN-1995  
CLASSIFICATION: 436  
ATTORNEY/AGENT INFORMATION:  
NAME: Mirock, S. Leslie  
REGISTRATION NUMBER: 18,872  
REFERENCE/DOCKET NUMBER: 1101-176  
TELECOMMUNICATION INFORMATION:  
TELEPHONE: (212) 790-9090  
TELEFAX: (212) 869-9741/8864  
TELEX: 66141 PENNIE  
INFORMATION FOR SEQ ID NO: 51:  
SEQUENCE CHARACTERISTICS:  
LENGTH: 43 amino acids  
TYPE: amino acid  
STRANDEDNESS: single  
TOPOLOGY: linear  
MOLECULE TYPE: peptide  
US-08-488-161-51  
Query Match 23.2%; Score 46; DB 2; Length 43;  
Best Local Similarity 43.8%; Pred. No. 4.9;  
Matches 7; Conservative 2; Mismatches 7; Indels 0; Gaps 0;  
QY 14 DPMKGPXKLMKGBG 29  
|:|:|:|:|  
Db 8 DPMKGPXKLMKGBG 23

Search completed: May 6, 2003, 14:59:00  
Job time : 16 secs



GenCore version 5.1.4 p5 4578  
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OM protein - protein search, using sw model

Run on: May 6, 2003, 14:54:59 ; Search time 34 Seconds  
(without alignments)  
141.089 Million cell updates/sec

Title: US-09-868-399-1  
Perfect score: 198  
Sequence: 1 KIQNFVRYRDSRDLPMKGPALKLMKGEAVIQDN 36

Scoring table: BLOSUM62  
Gapop 10.0 , Gapext 0.5

Searched: 908470 seqs, 133250620 residues

Total number of hits satisfying chosen parameters: 575469

Minimum DB seq length: 0  
Maximum DB seq length: 100

Post-processing: Minimum Match 0%  
Maximum Match 100%  
Listing first 45 summaries

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23: /SIDS2/gcgdata/geneSeq/geneSeq-emb1/AA2002.DAT.\*

Pred. No. is the number of results predicted by chance to have a score greater than or equal to the score of the result being printed, and is derived by analysis of the total score distribution.

#### SUMMARIES

Result No.	Score	Query Match	Length	DB	ID	Description
1	198	100.0	36	21	AA15339	HIV integrase pept
2	169	85.4	31	9	AA182730	Immunologically mi
3	169	85.4	50	9	AA182728	Immunologically mi
4	161	81.3	30	23	AA184494	HIV POL segment 63
5	110	55.6	30	23	AA184493	HIV POL segment 62
6	90	45.5	15	22	ABP24758	HIV DR super motif
7	88	44.4	15	22	ABP24701	HIV DR super motif
8	88	44.4	15	22	ABP25020	HIV DR 3b motif po
9	86	43.4	15	22	ABP24677	HIV DR super motif
10	86	43.4	15	22	ABP25016	HIV DR 3b motif po

11	83	41.9	30	23	AA184495	HIV POL segment 64
12	79	39.9	15	29	AA185346	Helper T-cell clas
13	79	39.9	15	22	ABP24643	HIV DR super motif
14	70	35.4	15	11	AA1803899	HIV antibody react
15	69	34.8	15	22	ABP24663	HIV DR super motif
16	68	34.3	11	22	ABP19154	HIV B62 super moti
17	66	33.3	11	22	ABP19309	HIV B62 super moti
18	62	31.3	10	22	ABP16223	HIV A24 super moti
19	62	31.3	10	22	ABP24215	HIV A24 motif pol
20	61	30.8	10	22	ABP16227	HIV A24 super moti
21	61	30.8	10	22	ABP21162	HIV A03 motif pol
22	61	30.8	10	22	ABP24230	HIV A24 motif pol
23	61	30.8	11	22	ABP13975	HIV A02 super moti
24	61	30.8	10	22	ABP16925	HIV B07 super moti
25	60	30.3	10	22	ABP16222	HIV A03 super moti
26	60	30.3	10	22	ABP21190	HIV A03 motif pol
27	60	30.3	10	22	ABP23194	HIV A11 motif pol
28	60	30.3	10	22	ABP24264	HIV A02 super moti
29	60	30.3	11	22	ABP13974	HIV A02 super moti
30	59	29.8	10	22	ABP16226	HIV A02 super moti
31	59	29.8	10	22	ABP16893	HIV B07 super moti
32	58	29.3	11	22	ABP23424	HIV A11 motif pol
33	58	29.3	11	22	ABP13973	HIV A02 super moti
34	58	29.3	11	22	ABP13976	HIV A02 super moti
35	58	29.3	11	22	ABP16382	HIV A24 super moti
36	58	29.3	11	22	ABP19302	HIV B62 super moti
37	58	29.3	11	22	ABP19464	E88, monoclonal an
38	58	29.3	43	17	AA195500	HIV B27 super moti
39	58	29.3	9	22	ABP17386	HIV A01 super moti
40	58	29.3	10	22	ABP11884	HIV A02 super moti
41	58	29.3	10	22	ABP13740	
42	57	28.8				
43	57	28.8				
44	57	28.8				
45	57	28.8				

#### ALIGNMENTS

RESULT 1  
AA15339  
ID AA15339 standard; peptide; 36 AA.  
AC AA15339;  
DT 15-DEC-2000 (first entry)  
XX  
DE HIV integrase peptide epitope.  
KW Detection; infection; HIV-1; antigen; pol; mixotrope; epitope; integrase;  
KW combinatorial peptide; antibody.  
XX  
OS Human immunodeficiency virus type 1.  
XX  
PN WO200040970-A1.  
PD 13-JUL-2000.  
XX  
PF 29-DEC-1999; 99WO-FR03311.  
XX  
PR 31-DEC-1998; 98FR-0016727.  
XX  
PA (INSP ) INST PASTEUR LILLE.  
PA (CNRS ) CNRS CENT NAT RECH SCI.  
XX (GRAS/) GRAS-MASSSE H.  
XX  
PI Tranchand-Bunel D, Aurault C;  
XX WPI; 2000-475858/41.  
DR  
XX  
PT Reagent for detecting human immune deficiency virus infection, useful  
for diagnosis and monitoring; comprises pol-derived antigen and mixture

PT of its convergent peptides -  
 XX  
 PS Claim 3; Page 15; 32pp; French.  
 XX  
 CC The invention relates to a reagent for detecting infection by human  
 CC immunodeficiency virus (HIV), comprising a mixture of an antigenic  
 CC peptide encoded by the pol gene and containing at most 60 (preferably  
 CC 20-40) amino acids, and a mixture, designated a mixotope, of converged,  
 CC combinatorial peptides derived from the antigenic peptide. This sequence  
 CC corresponds to an epitope from the integrase protein encoded by the pol  
 CC gene from HIV-1. The reagent is used to detect anti-HIV antibodies,  
 CC specifically for diagnosis and monitoring of infection.  
 XX  
 SQ Sequence 36 AA;  
 Query Match 100.0%; Score 198; DB 21; Length 36;  
 Best Local Similarity 100.0%; Pred. No. 2.8e-22;  
 Matches 36; Conservative 0; Mismatches 0; Indels 0; Gaps 0;  
 QY 1 KIQNFRVYRDSRDP LMKGPATLWKGEAVYIQDN 36  
 Db 1 KIQNFRVYRDSRDP LMKGPATLWKGEAVYIQDN 36  
 RESULT 2  
 AAP82730  
 ID AAP82730 standard; protein; 31 AA.  
 XX  
 AC AAP82730;  
 XX  
 DT 23-NOV-1990 (first entry)  
 XX  
 DE Immunologically mimicking peptide.  
 XX  
 KW Immunological mimicry; HIV gag and pol; AIDS.  
 XX  
 OS synthetic.  
 XX  
 PN EP267802-A.  
 XX  
 PD 18-MAY-1988.  
 XX  
 PF 13-NOV-1987; 87EP-0310047.  
 XX  
 PR 14-NOV-1986; 86US-0930785.  
 XX  
 PA (GENE-) GENETIC SYST CORP.  
 XX  
 PI Cosand WL, Harris LJ, Houghton RL;  
 XX  
 DR WPI; 1988-134645/20.  
 DR N-PSDB; AAN82239.  
 XX  
 PT Synthetic peptide(s) having sequence corresp. to HIV virus - used  
 PT for detection of AIDS-related disease and in vaccine prepn.  
 XX  
 PS Claim 14; Page 12; 12pp; English.  
 XX  
 CC This peptide immunologically mimics a protein encoded by the HIV  
 CC retrovirus and corresponds to part of peptide 123. It is competitive  
 CC with HIV and complexes formed on binding of antibodies are detected.  
 CC It is therefore used as an accurate test for detecting patients who  
 CC have been exposed to the aetiological agent of lymphadenopathy  
 CC syndrome and/or AIDS. Residue 31 can be absent but is opt. present.  
 CC See also AAN82236-38.  
 CC  
 XX  
 SQ Sequence 31 AA;  
 Query Match 85.4%; Score 169; DB 9; Length 31;  
 Best Local Similarity 100.0%; Pred. No. 5e-18;  
 Matches 30; Conservative 0; Mismatches 0; Indels 0; Gaps 0;  
 QY 1 KIQNFRVYRDSRDP LMKGPATLWKGEA 30

Db 1 KIQNFRVYRDSRDP LMKGPATLWKGEA 30  
 RESULT 3  
 AAP82728  
 ID AAP82728 standard; protein; 50 AA.  
 XX  
 AC AAP82728;  
 XX  
 DT 23-NOV-1990 (first entry)  
 XX  
 DE Immunologically mimicking peptide.  
 XX  
 KW Immunological mimicry; HIV gag and pol; AIDS.  
 XX  
 OS synthetic.  
 XX  
 PN EP267802-A.  
 XX  
 PD 18-MAY-1988.  
 XX  
 PF 13-NOV-1987; 87EP-0310047.  
 XX  
 PR 14-NOV-1986; 86US-0930785.  
 XX  
 PA (GENE-) GENETIC SYST CORP.  
 XX  
 PI Cosand WL, Harris LJ, Houghton RL;  
 XX  
 DR WPI; 1988-134645/20.  
 DR N-PSDB; AAN82237.  
 XX  
 PT Synthetic peptide(s) having sequence corresp. to HIV virus - used  
 PT for detection of AIDS-related disease and in vaccine prepn.  
 XX  
 PS Claim 14; Page 12; 12pp; English.  
 XX  
 CC This peptide which immunologically mimics a protein encoded by the  
 CC HIV retrovirus corresponds to amino acids 4385 to 4519 of pol  
 CC protein p31. It is competitive with HIV and complexes formed on  
 CC binding of antibodies are detected. It is therefore used as an  
 CC accurate test for detecting patients who have been exposed to the  
 CC aetiological agent of lymphadenopathy syndrome and/or AIDS.  
 CC Residues 1-4 and 50 can be absent but are opt. present.  
 CC See also AAN82236 and AAN82238-39.  
 CC  
 XX  
 SQ Sequence 50 AA;  
 Query Match 85.4%; Score 169; DB 9; Length 50;  
 Best Local Similarity 100.0%; Pred. No. 8.9e-18;  
 Matches 30; Conservative 0; Mismatches 0; Indels 0; Gaps 0;  
 QY 1 KIQNFRVYRDSRDP LMKGPATLWKGEA 30  
 Db 20 KIQNFRVYRDSRDP LMKGPATLWKGEA 49  
 RESULT 4  
 AAN84494  
 ID AAN84494 standard; Peptide; 30 AA.  
 XX  
 AC AAN84494;  
 XX  
 DT 08-MAY-2002 (first entry)  
 XX  
 DE HIV POL segment 63.  
 XX  
 KW Savine; vaccine; cancer; viral infection; HIV; hepatitis C virus;  
 KW viral infection; human immunodeficiency virus; melanoma;  
 KW bacterial infection; Salmonella; Legionella; parasitic infection;  
 KW Trypanosoma; Toxoplasma; Giardia.  
 XX

OS	Human immunodeficiency virus type 1.
OS	Synthetic.
XX	
XX	WO200190197-A1.
XX	
XX	29-NOV-2001.
XX	
XX	25-MAY-2001; 2001WO-AU00622.
XX	
XX	26-MAY-2000; 2000AU-0007761.
XX	
XX	(AUSU ) UNIV AUSTRALIAN NAT.
XX	
XX	Thomson SA, Ramshaw IA;
XX	
XX	WPI; 2002-147575/19.
XX	
XX	N-PSDB; ABK36333.
XX	
XX	New synthetic polypeptides having several different segments of at
XX	least one parent polypeptide linked together differently compared to
XX	the linkage in the parent polypeptide, for inducing immune response
XX	against a pathogen or cancer
XX	
XX	Example 1; Fig 12; 364pp; English.
XX	
XX	The invention relates to a new synthetic polypeptide (I) comprising
XX	several different segments of at least one parent polypeptide linked
XX	together in a different relationship relative to their linkage in the
XX	parent polypeptide to impede, abrogate or otherwise alter at least one
XX	function associated with the parent polypeptide and for inducing an
XX	immune response against a pathogen or cancer. Also included are a
XX	synthetic polynucleotide encoding and a computer system for
XX	designating the synthetic polypeptides. The synthetic polypeptides and
XX	polynucleotides are referred to as a Savine. The synthetic polypeptide is
XX	useful for modulating immune responses preferably directed against a
XX	pathogen or a cancer. (e.g., cancers of the lung, breast, ovary, cervix,
XX	colon, head and neck, pancreas, prostate, stomach, bladder, kidney, bone
XX	liver, oesophagus, brain, testicle, uterus), as potentiating agents.
XX	Compositions comprising the polypeptide may be used in the treatment or
XX	prophylaxis against viral (such as infections caused by HIV (human
XX	immunodeficiency virus), hepatitis, influenza, Japanese encephalitis
XX	virus, Epstein-Barr virus and respiratory syncytial virus), bacterial
XX	(e.g., infections caused by Neisseria, Meningococcal, Haemophilus,
XX	Salmonella, Streptococcal, Legionella and Mycobacterium) or parasitic
XX	(e.g., infections caused by Plasmodium, Schistosoma, Leishmania,
XX	Trypanosoma, Toxoplasma and Giardia) infections. The present
XX	sequence is a peptide derived from a parent protein used to
XX	construct a savine of the invention.
XX	
XX	Sequence 30 AA;
XX	
XX	Query Match 81.3%; Score 161; DB 23; Length 30;
XX	Best Local Similarity 96.7%; Pred No. 7, 5e-17;
XX	Matches 29; Conservative 0; Mismatches 1; Indels 0; Gaps 0
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XX	6 RYVYRDSRDLWKKGPAKILMKKGAVVIO 35
XX	
XX	1 RYVYRDSRDPXWKGPAPKILMKKGAVVIO 30
XX	
XX	
XX	RESULT 5
XX	AAU84493
XX	AAU84493 standard; Peptide; 30 AA.
XX	
XX	AAU84493;
XX	
XX	08-MAY-2002 (first entry)
XX	
XX	HIV POL segment 62.
XX	
XX	Savine; vaccine; cancer; viral infection; HIV; hepatitis C virus;
XX	viral infection; human immunodeficiency virus; melanoma;
XX	bacterial infection; Salmonella; Legionella; parasitic infection;

```

KW Trypanosoma; Toxoplasma; Giardia.
XX
XX Human immunodeficiency virus type 1.
OS Synthetic.
XX
XX WO200190197-A1.
XX
XX 29-NOV-2001.
XX
XX 25-MAY-2001; 2001WO-AU00622.
XX
XX 26-MAY-2000; 2000AU-000761.
XX
XX (AUSU ) UNIV AUSTRALIAN NAT.
XX
XX Thomson SA, Ramshaw IA;
XX
XX MPI; 2002-147575/19.
XX
XX N-PDB; ABK36332.
XX
XX
XX New synthetic polypeptides having several different segments of at
XX least one parent polypeptide linked together differently compared to
XX the linkage in the parent polypeptide, for inducing immune response
XX against a pathogen or cancer
XX
XX Example 1; Fig 12; 364pp; English.
XX
XX The invention relates to a new synthetic polypeptide (I) comprising
XX several different segments of at least one parent polypeptide linked
XX together in a different relationship relative to their linkage in the
XX parent polypeptide to impede, abrogate or otherwise alter at least one
XX function associated with the parent polypeptide and for inducing an
XX immune response against a pathogen or cancer. Also included are a
XX synthetic polynucleotide encoding and a computer system for
XX designing the synthetic polypeptides. The synthetic polypeptides and
XX polynucleotides are referred to as a Savine. The synthetic polypeptide is
XX useful for modulating immune responses preferably directed against a
XX pathogen or a cancer, (e.g., cancers of the lung, breast, ovary, cervix,
XX colon, head and neck, pancreas, prostate, stomach, bladder, kidney, bone
XX liver, oesophagus, brain, testicle, uterus), as potentiating agents.
XX Compositions comprising the polypeptide may be used in the treatment or
XX prophylaxis against viral (such as infections caused by HIV (human
XX immunodeficiency virus), hepatitis, influenza, Japanese encephalitis
XX virus, Epstein-Barr virus and respiratory syncytial virus), bacterial
XX (e.g., infections caused by Neisseria, Meningococcal, Haemophilus,
XX Salmonella, Streptococcal, Legionella and Mycobacterium or parasitic
XX (e.g., infections caused by Plasmodium, Schistosoma, Leishmania,
XX Trypanosoma, Toxoplasma and Giardia) infections. The present
XX sequence is a peptide derived from a parent protein used to
XX construct a savine of the invention.
XX
XX Sequence 30 AA;
XX
XX Query Match 55.6%; Score 110; DB 23; Length 30;
XX Best Local Similarity 95.0%; Pred. No. 3.1e-09;
XX Matches 19; Conservative 0; Mismatches 1; Indels 0; Gaps 0;
XX
XX 1 KIQNFRVYRRDSRDLWKGP 20
XX |||||
XX 11 KIQNFRVYRRDSRDLWKGP 30
XX
XX RESULT 6
XX ABP24758
XX ID ABP24758 standard; Peptide: 15 AA.
XX AC ABP24758;
XX XX
XX DT 15-JUL-2002 (first entry)
XX DE HIV DR super motif pol peptide #125.
XX
XX HIV, HIV-1; human immunodeficiency virus; env; pol; gag; nef; vpr;
XX

```

KW vpu; vif; tat; cytotoxic T lymphocyte; CTL; immune response; epitope;  
 KM antigen; vaccine; HIV infection; immunisation; virucide.  
 OS Human immunodeficiency virus type 1.  
 XX WO200124810-A1.  
 XX PN 12-APR-2001.  
 XX PD 05-OCT-2000; 2000WO-US27766.  
 XX PF 05-OCT-1999; 99US-0412863.  
 XX PR (EPIM-) EPIMUNE INC.  
 XX PI Sette A, Sidney J, Southwood S, Livingston BD, Chesnut R;  
 XX PI Baker DM, Celis E, Kubo RT, Grey HM,  
 XX DR WPI; 2001-354887/37.  
 XX PT Vaccine compositions comprising human immunodeficiency virus-1 (HIV-1)  
 XX PT peptide groups, useful for vaccinating against HIV-1 -  
 XX PS Claim 32; Page 376; 448pp; English.  
 XX CC The present invention describes a composition (I) comprising a prepared  
 CC human immunodeficiency virus-1 (HIV-1) group comprising an amino acid  
 CC sequence selected from 51 defined amino acid sequences (AB25347 to  
 CC AB25397). (I) has virucide activity and can be used in vaccines. (I)  
 CC may be used for immunising subjects against HIV-1 infections. The use of  
 CC group-based vaccines has several advantages over traditional vaccines,  
 CC particularly when compared to the use of whole antigens in vaccine  
 CC compositions. There is evidence that the immune response to whole  
 CC antigens is directed largely toward variable regions of the antigen,  
 CC allowing for immune escape due to mutations. The groups for inclusion in  
 CC an group-based vaccine may be selected from conserved regions of viral or  
 CC tumour-associated antigens, which therefore reduces the likelihood of  
 CC escape mutants. Furthermore, immunosuppressive groups that may be present  
 CC in whole antigens can be avoided with the use of group-based vaccines.  
 CC An additional advantage of an group-based vaccine approach is the ability  
 CC to combine selected groups (CTL and HTL), and further, to modify the  
 CC composition of the groups, achieving, for example, enhanced  
 CC immunogenicity. Accordingly, the immune response can be modulated, as  
 CC appropriate, for the target disease. Similar engineering of the response  
 CC is not possible with traditional approaches. ABP1501 to ABP25412  
 CC represent peptide sequences used in the exemplification of the present  
 CC invention.  
 CC SQ Sequence 15 AA;  
 CC  
 CC Query Match 45.5%; Score 90; DB 22; Length 15;  
 CC Best Local Similarity 100.0%; Pred. No. 1.3e-06;  
 CC Matches 15; Conservative 0; Mismatches 0; Indels 0; Gaps 0;  
 CC  
 CC QY 13 RDPLMKGPATKLWKG 27  
 CC |||||  
 CC Db 1 RDPLMKGPATKLWKG 15  
 CC  
 CC RESULT 7  
 CC ID ABP24701 standard; Peptide; 15 AA.  
 CC AC ABP24701;  
 CC XX  
 CC DT 15-JUL-2002 (first entry)  
 CC XX  
 CC DE HIV DR super motif pol peptide #68.  
 CC XX  
 CC KM HIV; HIV-1; human immunodeficiency virus; env; pol; gag; nef; vpr;  
 CC KM vpu; vif; tat; cytotoxic T lymphocyte; CTL; immune response; epitope;  
 CC KM antigen; vaccine; HIV infection; immunisation; virucide.  
 CC XX

OS Human immunodeficiency virus type 1.  
 XX WO200124810-A1.  
 XX PN 12-APR-2001.  
 XX PD 05-OCT-2000; 2000WO-US27766.  
 XX PF 05-OCT-1999; 99US-0412863.  
 XX PR (EPIM-) EPIMUNE INC.  
 XX PI Sette A, Sidney J, Southwood S, Livingston BD, Chesnut R;  
 XX PI Baker DM, Celis E, Kubo RT, Grey HM,  
 XX DR WPI; 2001-354887/37.  
 XX PT Vaccine compositions comprising human immunodeficiency virus-1 (HIV-1)  
 XX PT peptide groups, useful for vaccinating against HIV-1 -  
 XX PS Claim 32; Page 375; 448pp; English.  
 XX CC The present invention describes a composition (I) comprising a prepared  
 CC human immunodeficiency virus-1 (HIV-1) group comprising an amino acid  
 CC sequence selected from 51 defined amino acid sequences (AB25347 to  
 CC ABP25397). (I) has virucide activity and can be used in vaccines. (I)  
 CC may be used for immunising subjects against HIV-1 infections. The use of  
 CC group-based vaccines has several advantages over traditional vaccines,  
 CC particularly when compared to the use of whole antigens in vaccine  
 CC compositions. There is evidence that the immune response to whole  
 CC antigens is directed largely toward variable regions of the antigen,  
 CC allowing for immune escape due to mutations. The groups for inclusion in  
 CC an group-based vaccine may be selected from conserved regions of viral or  
 CC tumour-associated antigens, which therefore reduces the likelihood of  
 CC escape mutants. Furthermore, immunosuppressive groups that may be present  
 CC in whole antigens can be avoided with the use of group-based vaccines.  
 CC An additional advantage of an group-based vaccine approach is the ability  
 CC to combine selected groups (CTL and HTL), and further, to modify the  
 CC composition of the groups, achieving, for example, enhanced  
 CC immunogenicity. Accordingly, the immune response can be modulated, as  
 CC appropriate, for the target disease. Similar engineering of the response  
 CC is not possible with traditional approaches. ABP1501 to ABP25412  
 CC represent peptide sequences used in the exemplification of the present  
 CC invention.  
 CC SQ Sequence 15 AA;  
 CC  
 CC Query Match 44.4%; Score 88; DB 22; Length 15;  
 CC Best Local Similarity 93.3%; Pred. No. 2.6e-06;  
 CC Matches 14; Conservative 1; Mismatches 0; Indels 0; Gaps 0;  
 CC  
 CC QY 13 RDPLMKGPATKLWKG 27  
 CC |||||  
 CC Db 1 RDPLMKGPATKLWKG 15  
 CC  
 CC RESULT 8  
 CC ID ABP25020 standard; Peptide; 15 AA.  
 CC AC ABP25020;  
 CC XX  
 CC DT 15-JUL-2002 (first entry)  
 CC XX  
 CC DE HIV DR 3b motif pol peptide #11.  
 CC XX  
 CC KM HIV; HIV-1; human immunodeficiency virus; env; pol; gag; nef; vpr;  
 CC KM vpu; vif; tat; cytotoxic T lymphocyte; CTL; immune response; epitope;  
 CC KM antigen; vaccine; HIV infection; immunisation; virucide.  
 CC OS Human immunodeficiency virus type 1.  
 CC XX  
 CC PN WO200124810-A1.

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XX 12-APR-2001.
XX
XX 05-OCT-2000; 2000WO-US27766.
XX
XX 05-OCT-1999; 99US-0412863.
XX
XX (EPIM-) EPIMMUNE INC.
XX
XX Sette A, Sidney J, Southwood S, Livingston BD, Chesnut R;
XX Baker DM, Cells E, Kubo RT, Grey HM;
XX
XX WPI; 2001-354887/37.
XX
XX Vaccine compositions comprising human immunodeficiency virus-1 (HIV-1)
XX peptide groups, useful for vaccinating against HIV-1 -
XX
XX Claim 32; Page 408; 448pp; English.
XX
XX The present invention describes a composition (I) comprising a prepared
XX human immunodeficiency virus-1 (HIV-1) group comprising an amino acid
XX sequence selected from 51 defined amino acid sequences (AB125347 to
XX ABP25397). (I) has virucide activity and can be used in vaccines. (I)
XX may be used for immunising subjects against HIV-1 infections. The use of
XX group-based vaccines has several advantages over traditional vaccines,
XX particularly when compared to the use of whole antigens in vaccine
XX compositions. There is evidence that the immune response to whole
XX antigens is directed largely toward variable regions of the antigen,
XX allowing for immune escape due to mutations. The groups for inclusion in
XX an group-based vaccine may be selected from conserved regions of viral or
XX tumour-associated antigens, which therefore reduces the likelihood of
XX escape mutants. Furthermore, immunosuppressive groups that may be present
XX in whole antigens can be avoided with the use of group-based vaccines.
XX An additional advantage of an group-based vaccine approach is the ability
XX to combine selected groups (CTL and HTL), and further, to modify the
XX composition of the groups, achieving, for example, enhanced
XX immunogenicity. Accordingly, the immune response can be modulated, as
XX appropriate, for the target disease. Similar engineering of the response
XX is not possible with traditional approaches. ABP11501 to ABP25412
XX represent peptide sequences used in the exemplification of the present
XX invention.
XX
XX Sequence 15 AA;
XX
XX Query Match 44.4%; Score 86; DB 22; Length 15;
XX Best Local Similarity 100.0%; Pred. No. 2.6e-06;
XX Matches 15; Conservative 0; Mismatches 0; Indels 0; Gaps 0;
XX
XX QY 5 FRVYRDSRDPIMKG 19
XX |||||
XX 1 FRVYRDSRDPIMKG 15
XX
XX Db
XX
XX RESULT 9
XX ABP24677
XX ID ABP24677 standard; Peptide; 15 AA.
XX
XX AC ABP24677;
XX
XX DT 15-JUL-2002 (first entry)
XX
XX DE HIV DR super motif pol peptide #44.
XX
XX KW HIV; HIV-1; human immunodeficiency virus; env; pol; gag; nef; vpr;
XX vpu; vif; tat; cytototoxic T lymphocyte; CTL; immune response; epitope;
XX antigen; vaccine; HIV infection; immunisation; virucide.
XX
XX OS Human immunodeficiency virus type 1.
XX
XX PN WO200124810-A1.
XX
XX PD 12-APR-2001.
XX

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PF 05-OCT-2000; 2000WO-US27766.
XX
XX 05-OCT-1999; 99US-0412863.
XX
XX (EPIM-) EPIMMUNE INC.
XX
XX Sette A, Sidney J, Southwood S, Livingston BD, Chesnut R;
XX Baker DM, Cells E, Kubo RT, Grey HM;
XX
XX WPI; 2001-354887/37.
XX
XX Vaccine compositions comprising human immunodeficiency virus-1 (HIV-1)
XX peptide groups, useful for vaccinating against HIV-1 -
XX
XX Claim 32; Page 374; 448pp; English.
XX
XX The present invention describes a composition (I) comprising a prepared
XX human immunodeficiency virus-1 (HIV-1) group comprising an amino acid
XX sequence selected from 51 defined amino acid sequences (AB125347 to
XX ABP25397). (I) has virucide activity and can be used in vaccines. (I)
XX may be used for immunising subjects against HIV-1 infections. The use of
XX group-based vaccines has several advantages over traditional vaccines,
XX particularly when compared to the use of whole antigens in vaccine
XX compositions. There is evidence that the immune response to whole
XX antigens is directed largely toward variable regions of the antigen,
XX allowing for immune escape due to mutations. The groups for inclusion in
XX an group-based vaccine may be selected from conserved regions of viral or
XX tumour-associated antigens, which therefore reduces the likelihood of
XX escape mutants. Furthermore, immunosuppressive groups that may be present
XX in whole antigens can be avoided with the use of group-based vaccines.
XX An additional advantage of an group-based vaccine approach is the ability
XX to combine selected groups (CTL and HTL), and further, to modify the
XX composition of the groups, achieving, for example, enhanced
XX immunogenicity. Accordingly, the immune response can be modulated, as
XX appropriate, for the target disease. Similar engineering of the response
XX is not possible with traditional approaches. ABP11501 to ABP25412
XX represent peptide sequences used in the exemplification of the present
XX invention.
XX
XX Sequence 15 AA;
XX
XX Query Match 43.4%; Score 86; DB 22; Length 15;
XX Best Local Similarity 93.3%; Pred. No. 5.2e-06;
XX Matches 14; Conservative 1; Mismatches 0; Indels 0; Gaps 0;
XX
XX QY 4 NFRVYRDSRDPIMK 18
XX |||||
XX 1 NFRVYRDSRDPIMK 15
XX
XX Db
XX
XX RESULT 10
XX ABP25016
XX ID ABP25016 standard; Peptide; 15 AA.
XX
XX AC ABP25016;
XX
XX DT 15-JUL-2002 (first entry)
XX
XX DE HIV DR 3b motif pol peptide #7.
XX
XX KW HIV; HIV-1; human immunodeficiency virus; env; pol; gag; nef; vpr;
XX vpu; vif; tat; cytototoxic T lymphocyte; CTL; immune response; epitope;
XX antigen; vaccine; HIV infection; immunisation; virucide.
XX
XX OS Human immunodeficiency virus type 1.
XX
XX PN WO200124810-A1.
XX
XX PD 12-APR-2001.
XX
XX PF 05-OCT-2000; 2000WO-US27766.
XX
XX PR 05-OCT-1999; 99US-0412863.
XX

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XX (EPTM-) EPIMUNE INC.  
XX

XX Sette A, Sidney J, Southwood S, Livingston BD, Chesnut R;  
XX Baker DM, Celis E, Kubo RT, Grey HM;  
XX WPI: 2001-35487/37.

XX Vaccine compositions comprising human immunodeficiency virus-1 (HIV-1)  
XX peptide groups, useful for vaccinating against HIV-1  
XX

PS Claim 32; Page 408; 448pp; English.

CC The present invention describes a composition (I) comprising a prepared  
CC human immunodeficiency virus-1 (HIV-1) group comprising an amino acid  
CC sequence selected from 51 defined amino acid sequences (AB125347 to  
CC AB25397). (I) has virucide activity and can be used in vaccines. (I)  
CC may be used for immunising subjects against HIV-1 infections. The use of  
CC group-based vaccines has several advantages over traditional vaccines,  
CC particularly when compared to the use of whole antigens in vaccine  
CC compositions. There is evidence that the immune response to whole  
CC antigens is directed largely toward variable regions of the antigen,  
CC allowing for immune escape due to mutations. The groups for inclusion in  
CC an group-based vaccine may be selected from conserved regions of viral or  
CC tumour-associated antigens, which therefore reduces the likelihood of  
CC in whole antigens can be avoided with the use of group-based vaccines.  
CC An additional advantage of an group-based vaccine approach is the ability  
CC to combine selected groups (CTL and HTL), and further, to modify the  
CC composition of the groups, achieving, for example, enhanced  
CC immunogenicity. Accordingly, the immune response can be modulated, as  
CC appropriate, for the target disease. Similar engineering of the response  
CC is not possible with traditional approaches. AB11501 to AB25412  
CC represent peptide sequences used in the exemplification of the present  
CC invention.

CC Sequence 15 AA:

Query Match Best Local Similarity 43.4%; Score 86; DB 22; Length 15;  
Matches 14; Conservative 1; Mismatches 0; Indels 0; Gaps 0;

OY 5 FRVYRDSRDLPMKG 19  
DB 1 FRVYRDSRDLPMKG 15

RESULT 11

AAU84495 standard; Peptide; 30 AA.

AC AAU84495;  
DT 08-MAY-2002 (first entry)  
XX  
DE HIV POL segment 64.

KW Savine; vaccine; cancer; viral infection; HIV; hepatitis C virus;  
KW viral infection; human immunodeficiency virus; melanoma;  
KW bacterial infection; Salmonella; Legionella; parasitic infection;  
KW Trypanosoma; Toxoplasma; Giardia.

OS Human immunodeficiency virus type 1.  
OS Synthetic.

PN WO200190197-A1.

PD 29-NOV-2001.

PF 25-MAY-2001; 2001WO-AU00622.

PR 26-MAY-2000; 2000AU-0007761.

PA (AUSU) UNIV AUSTRALIAN NAT.

XX Thomson SA, Ramsdew IA;  
XX

DR WPI: 2002-147575/19.  
DR N-PSDB; ABK36334.

PT New synthetic polypeptides having several different segments of at  
PT least one parent polypeptide linked together differently compared to  
PT the linkage in the parent polypeptide, for inducing immune response  
PT against a pathogen or cancer  
XX

PS Example 1; Fig 12; 364pp; English.

CC The invention relates to a new synthetic polypeptide (I) comprising  
CC several different segments of at least one parent polypeptide linked  
CC together in a different relationship relative to their linkage in the  
CC parent polypeptide to impede, abrogate or otherwise alter at least one  
CC function associated with the parent polypeptide and for inducing an  
CC immune response against a pathogen or cancer. Also included are a  
CC synthetic polynucleotide encoding and a computer system for  
CC designing the synthetic polypeptides. The synthetic polypeptides and  
CC polynucleotides are referred to as a Savine. The synthetic polypeptide is  
CC useful for modulating immune responses preferably directed against a  
CC pathogen or a cancer, (e.g., cancers of the lung, breast, ovary, cervix,  
CC colon, head and neck, pancreas, prostate, stomach, bladder, kidney, bone  
CC liver, oesophagus, brain, testicle, uterus), as potentiating agents.  
CC Compositions comprising the polypeptide may be used in the treatment or  
CC prophylaxis against viral (such as infections caused by HIV (human  
CC virus, Epstein-Barr virus and respiratory syncytial virus), bacterial  
CC (e.g., infections caused by Neisseria, Meningococcal, Haemophilus,  
CC Salmonella, Streptococcal, Legionella, and Mycobacterium) or parasitic  
CC (e.g., infections caused by Plasmodium, Schistosoma, Leishmania,  
CC Trypanosoma, Toxoplasma and Giardia) infections. The present  
CC sequence is a peptide derived from a parent protein used to  
CC construct a Savine of the invention.

CC Sequence 30 AA:

Query Match Best Local Similarity 41.9%; Score 83; DB 23; Length 30;  
Matches 16; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

OY 21 AKLWKGAGVAVIODN 36  
DB 1 AKLWKGAGVAVIODN 16

RESULT 12

AAW85346 standard; peptide; 15 AA.

AC AAW85346;

DT 16-FEB-1999 (first entry)

DE Helper T-cell class II peptide derived from POL protein.

KW Helper T-cell peptide; human leucocyte antigen; HLA; DR4w4; DR1;  
KW DR7; cytotoxic T lymphocyte; CTL; hepatitis; autoimmune disease;  
KW acquired immune deficiency syndrome; malaria; cancer;  
KW allograft rejection; allergy; Lyme disease; hepatitis;  
KW post-streptococcal endocarditis; glomerulonephritis;  
KW food hypersensitivity.

OS Synthetic.

OS Human immunodeficiency virus type 1.

PN WO9832456-A1.

PD 30-JUL-1998.

PF 23-JAN-1998; 98WO-US01373.  
XX  
PT 07-FEB-1997; 97US-0037432.  
PR 23-JAN-1997; 97US-0036713.  
XX  
PA (EPIM-) EPIMMUNE INC.  
XX  
PI Sette A, Sidney J, Southwood S;  
XX  
DR WPI; 1998-427679/36.  
XX  
PT Composition containing peptide that induces cytotoxic T lymphocyte  
PT response, and helper peptide - can bind to human leucocyte antigen  
PT alleles, used to treat or prevent cancers, parasitic infections and  
PT autoimmune disease  
XX  
PS Disclosure; Page 41; 51pp; English.  
XX  
XX AAM85284-451 represent helper T-cell class II peptides, which can bind  
CC to the human leucocyte antigens (HLA) DR4w4, DRI and DR7. The peptides  
CC are used in the course of the invention. The specification describes  
CC peptides that induce a cytotoxic T lymphocyte (CTL) response, and  
CC T-helper peptides, that are used together to generate a CTL response for  
CC the treatment or prevention of viral, fungal, bacterial or parasitic  
CC infections (e.g. hepatitis, acquired immune deficiency syndrome or  
CC malaria) or cancer (e.g. renal or cervical carcinoma, lymphoma, prostate  
CC cancer or condyloma acuminatum). Helper T-cell peptides may be used  
CC alone to induce a helper T cell response, e.g. in cases of autoimmune  
CC disease, allograft rejection, allergy, Lyme disease, hepatitis,  
CC post-streptococcal endocarditis, glomerulonephritis and food  
CC hypersensitivity.  
XX  
SQ Sequence 15 AA;  
XX  
XX  
Query Match 39.9%; Score 79; DB 19; Length 15;  
Best Local Similarity 100.0%; Pred. No. 5.7e-05;  
Matches 15; Conservative 0; Mismatches 0; Indels 0; Gaps 0;  
XX  
QY 22 KILMKSGAVVTDN 36  
DB 1 KILMKSGAVVTDN 15  
XX  
XX  
RESULT 13  
ID AABP24643 standard; Peptide; 15 AA.  
XX  
AC AABP24643;  
XX  
DT 15-JUL-2002 (first entry)  
XX  
DE HIV DR super motif pol peptide #10.  
XX  
XX HIV; HIV-1; human immunodeficiency virus; env; pol; gag; nef; vpr;  
XX vpu; vif; tat; cytototoxic T lymphocyte; CTL; immune response; epitope;  
XX antigen; vaccine; HIV infection; immunisation; virucide.  
XX  
OS Human immunodeficiency virus type 1.  
XX  
XX WO200124810-A1.  
XX  
XX 12-APR-2001.  
XX  
XX 05-OCT-2000; 2000WO-US27766.  
XX  
XX 05-OCT-1999; 99US-0412863.  
XX  
XX (EPIM-) EPIMMUNE INC.  
XX  
XX Sette A, Sidney J, Southwood S, Livingston BD, Chesnut R;  
XX Baker DM, Cells E, Kubo RT, Grey HM;  
XX  
XX WPI; 2001-354887/37.

XX  
XX Vaccine compositions comprising human immunodeficiency virus-1 (HIV-1)  
PT peptide groups, useful for vaccinating against HIV-1 -  
XX  
XX Claim 32; Page 374; 448pp; English.  
XX  
XX The present invention describes a composition (I) comprising a prepared  
CC human immunodeficiency virus-1 (HIV-1) group comprising an amino acid  
CC sequence selected from 51 defined amino acid sequences (AB125347 to  
CC ABP25397). (II) has virucide activity and can be used in vaccines. (I)  
CC may be used for immunising subjects against HIV-1 infections. The use of  
CC group-based vaccines has several advantages over traditional vaccines,  
CC particularly when compared to the use of whole antigens in vaccine  
CC compositions. There is evidence that the immune response to whole  
CC antigens is directed largely toward variable regions of the antigen,  
CC allowing for immune escape due to mutations. The groups for inclusion in  
CC an group-based vaccine may be selected from conserved regions of viral or  
CC tumour-associated antigens, which therefore reduces the likelihood of  
CC escape mutants. Furthermore, immunosuppressive groups that may be present  
CC in whole antigens can be avoided with the use of group-based vaccines.  
CC An additional advantage of an group-based vaccine approach is the ability  
CC to combine selected groups (CTL and HTL), and further, to modify the  
CC composition of the groups, achieving, for example, enhanced  
CC immunogenicity. Accordingly, the immune response can be modulated, as  
CC appropriate, for the target disease. Similar engineering of the response  
CC is not possible with traditional approaches. ABP11501 to ABP25412  
CC represent peptide sequences used in the exemplification of the present  
CC invention.  
XX  
SQ Sequence 15 AA;  
XX  
XX  
Query Match 39.9%; Score 79; DB 22; Length 15;  
Best Local Similarity 100.0%; Pred. No. 5.7e-05;  
Matches 15; Conservative 0; Mismatches 0; Indels 0; Gaps 0;  
XX  
QY 22 KILMKSGAVVTDN 36  
DB 1 KILMKSGAVVTDN 15  
XX  
XX  
RESULT 14  
ID AAR03899 standard; peptide; 15 AA.  
XX  
AC AAR03899;  
XX  
DT 16-FEB-1993 (first entry)  
XX  
DE HIV-antibody reactive peptide (2).  
XX  
XX HIV; diagnosis.  
XX  
XX Synthetic.  
XX  
XX EP362915-A.  
XX  
XX 11-APR-1990.  
XX  
XX 07-SEP-1989; 89EP-0202258.  
XX  
XX 09-SEP-1988; 88NL-0002217.  
XX  
XX (ALKU) AKZO NV.  
XX  
XX Hellings JA, Schalcken JJ, Sprengers ED;  
XX  
XX WPI; 1990-109271/15.  
XX  
XX  
PT New synthetic oligopeptide cpds. reactive with HIV antibodies -  
XX useful as diagnostic reagents are non infectious and safe to use  
XX  
XX Claim 3; Page 5 + Fig 2; 7pp; English.  
XX

CC The peptides given in AAR03898-900 react immunochemically with  
 CC antibodies directed against HIV. They are suitable for use in a  
 CC diagnostic method for determining the presence of HIV or HIV-  
 CC antibodies in a test fluid. In contrast to the native HIV, the  
 CC peptides have the great advantage that these are of a safe non-  
 CC infectious origin. They are made by standard (esp. solid phase)  
 CC methods of peptide synthesis, or by recombinant DNA techniques.

XX  
 SQ Sequence 15 AA;

Query Match

Best Local Similarity 35.4%; Score 70; DB 11; Length 15;  
 Matches 13; Conservative 1; Mismatches 0; Indels 0; Gaps 0;

OY 1 KIQNFRVYRDSRD 14  
 |||||  
 Db 2 KIQNFRVYRDSRN 15

RESULT 15

ABP24663  
 ID ABP24663 standard; Peptide; 15 AA.

XX ABP24663;

DT 15-JUL-2002 (first entry)

DE HIV DR super motif pol peptide #30.

KW HIV; HIV-1; human immunodeficiency virus; env; pol; gag; nef; vpr;  
 VPU; vif; tat; cytoxic T lymphocyte; CTL; immune response; epitope;  
 antigen; vaccine; HIV infection; immunisation; virucide.

OS Human immunodeficiency virus type 1.

FN WO200124810-A1.

PD 12-APR-2001.

PF 05-OCT-2000; 2000WO-US27766.

PR 05-OCT-1999; 99US-0412863.

PA (EPLM-) EPIMUNE INC.

PI Sette A, Sidney J, Southwood S, Livingston BD, Chesnut R;  
 PI Baker DM, Celis E, Kudo RT, Grey HM;

DR WPI; 2001-354887/37.

Vaccine compositions comprising human immunodeficiency virus-1 (HIV-1)  
 peptide groups, useful for vaccinating against HIV-1 -

Claim 32; Page 374; 448pp; English.

CC The present invention describes a composition (I) comprising a prepared  
 CC human immunodeficiency virus-1 (HIV-1) group comprising an amino acid  
 CC sequence selected from 51 defined amino acid sequences (ABU25347 to  
 CC ABP25397). (I) has virucide activity and can be used in vaccines. (I)  
 CC may be used for immunising subjects against HIV-1 infections. The use of  
 CC group-based vaccines has several advantages over traditional vaccines,  
 CC particularly when compared to the use of whole antigens in vaccine  
 CC compositions. There is evidence that the immune response to whole  
 CC antigens is directed largely toward variable regions of the antigen,  
 CC allowing for immune escape due to mutations. The groups for inclusion in  
 CC an group-based vaccine may be selected from conserved regions of viral or  
 CC tumour-associated antigens, which therefore reduces the likelihood of  
 CC escape mutants. Furthermore, immunosuppressive groups that may be present  
 CC in whole antigens can be avoided with the use of group-based vaccines.  
 CC An additional advantage of an group-based vaccine approach is the ability  
 CC to combine selected groups (CTL and HTL), and further, to modify the  
 CC composition of the groups, achieving, for example, enhanced  
 CC immunogenicity. Accordingly, the immune response can be modulated, as

CC appropriate, for the target disease. Similar engineering of the response  
 CC is not possible with traditional approaches. ABP11501 to ABP25412  
 CC represent peptide sequences used in the exemplification of the present  
 CC invention.

XX  
 SQ Sequence 15 AA;

Query Match

Best Local Similarity 34.8%; Score 69; DB 22; Length 15;  
 Matches 13; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

OY 1 KIQNFRVYRDSR 13  
 |||||  
 Db 3 KIQNFRVYRDSR 15

Search completed: May 6, 2003, 14:57:23  
 Job time : 35 secs